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OM protein - protein search, using sw model

Run on: March 20, 2000, 05:31:20 ; Search time 35.25 Seconds
(without alignments)
249.964 Million cell updates/sec

Title: US-08-509-359B-138
Perfect score: 1923
Sequence: 1 EELTKYGAHVIMLFVPVT.....STDNLVRPFMDTLASHQLYI 372

Scoring table: BLOSUM62

Searched: 188963 seqs, 23686106 residues

Database : A_Geneseq_36:*

Word size : 0

Number of hits that pass the threshold : 188963

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1923	100.0	448	1 W05762	Human presenilin-2
2	1923	100.0	448	1 W11321	Human AD4 protein.
3	1923	100.0	448	1 W23967	Human presenilin-2
4	1919	99.8	448	1 W05763	Presenilin-2 M239V
5	1918	99.7	448	1 W05765	Presenilin-2 I420T
6	1914	99.5	448	1 W05764	Presenilin-2 N141I
7	1907.5	99.2	447	1 W28508	Full AD4/AD3LP seq
8	1723	89.6	414	1 W05766	Presenilin-2 delta
9	1591.5	82.8	376	1 W28506	AD4/AD3LP sequence
10	1431.5	74.4	467	1 W05735	Murine presenilin.
11	1431.5	74.4	467	1 W23966	Mouse presenilin-1
12	1429.5	74.3	467	1 W11839	Human early onset
13	1429.5	74.3	463	1 W11840	Early onset Alzhei
14	1429.5	74.3	467	1 W05733	Presenilin-1-1. Ne
15	1429.5	74.3	407	1 W28507	Partial AD3 sequen
16	1429.5	74.3	467	1 W41430	PS1/467 protein. D
17	1429.5	74.3	429	1 W41429	PS1/429 protein. D
18	1429.5	74.3	467	1 W23964	Human presenilin-1
19	1426.5	74.2	467	1 W05755	Presenilin-1-1 L28
20	1426.5	74.2	467	1 W05758	Presenilin-1-1 L39
21	1426.5	74.2	467	1 W05737	Presenilin-1-1 V82
22	1426.5	74.2	467	1 W05746	Presenilin-1-1 I21
23	1426.5	74.2	467	1 W41431	Mouse PS1/467 prot
24	1425.5	74.1	467	1 W05754	Presenilin-1-1 A28
25	1425.5	74.1	467	1 W05736	Presenilin-1-1 A79
26	1425.5	74.1	467	1 W05747	Presenilin-1-1 I23
27	1425.5	74.1	467	1 W05749	Presenilin-1-1 A26
28	1424.5	74.1	463	1 W12376	Human S182 gene pr
29	1424.5	74.1	467	1 W05738	Presenilin-1-1 V96
30	1424.5	74.1	467	1 W05739	Presenilin-1-1 Y11
31	1424.5	74.1	467	1 W05741	Presenilin-1-1 I14
32	1424.5	74.1	467	1 W05748	Presenilin-1-1 A24
33	1424.5	74.1	467	1 W27176	Human S182 gene, P
34	1423.5	74.0	467	1 W05753	Presenilin-1-1 E28
35	1423.5	74.0	467	1 W05757	Presenilin-1-1 G38
36	1423.5	74.0	463	1 W05734	Presenilin-1-2. Ne
37	1423.5	74.0	467	1 W05740	Presenilin-1-1 M13
38	1423.5	74.0	467	1 W05742	Presenilin-1-1 M14
39	1423.5	74.0	463	1 W23965	Human presenilin-1

40 1422.5 74.0 467 1 W05744 Presenilin-1-1 L17
41 1422.5 74.0 467 1 W56770 Homo sapiens PS-1.
42 1421.5 73.9 467 1 W05752 Presenilin-1-1 P26
43 1421.5 73.9 463 1 W42375 Human presenilin 1
44 1420.5 73.9 467 1 W05743 Presenilin-1-1 H16
45 1420.5 73.9 467 1 W05745 Presenilin-1-1 G20

ALIGNMENTS

RESULT 1

W05762

ID W05762 standard; Protein; 448 AA.

AC W05762;

DT 25-JUL-1997 (first entry)

DE Human presenilin-2.

KW Presenilin-2; human; hPS1-1; hPS1-2; PS-2; integral membrane protein; AD;

KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;

KW depression; antibody; gene expression modulator; therapy.

OS Homo sapiens.

PN W09634099-A2.

PD 31-OCT-1996.

PF 29-APR-1996; CA0263.

PR 28-APR-1995; US-431048.

PR 28-JUN-1995; US-496841.

PR 31-JUL-1995; US-509359.

PA (HSCR-) HSC RES & DEV LP.

PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.

PI Fraser PE, Rommens JM, St George-Hyslop PH;

DR WPI: 96-497631/49.

DR N-PSDB; T40031.

PT New presenilin genes - useful for diagnosis, therapy and drug

PT screening of familial Alzheimer's disease, cerebral disorders, etc.

PS Claim 4; Page 148-150; 178pp; English.

CC This sequence represents the wild type human presenilin-2.

CC W05733 and W05734 represent the two different forms of wild type human

CC presenilin-1 (PS-1). The form represented by W05734 results from

CC alternate splicing of the genomic DNA sequence. W05735 represents the

CC coding sequence for wild type mouse PS-1. The presenilins are a family of

CC highly conserved integral membrane proteins with a common structural

CC motif, common alternate splicing patterns, and common mutational hot spot

CC regions. Mutations in PS genes are implicated in familial Alzheimer's

CC disease (AD) and possibly other diseases such as cerebral haemorrhage,

CC schizophrenia, depression etc., so detection of mutations in the DNA

CC encoding these sequences can be used for diagnosis of these diseases.

CC These proteins, or vectors that express them or containing antisense

CC sequences, antibodies selective for mutant forms of these proteins (such

CC as W05736) and modulators of PS gene expression are potentially useful

CC for treatment of AD etc. Transgenic animals are useful as models for drug

CC screening. The antibodies can also be used e.g. for affinity purification

CC and in immunoassays.

CC Sequence 448 AA;

SQ

Query Match 100.0%; Score 1923; DB 1; Length 448;

Best Local Similarity 100.0%; Pred. No. 1.6e-200;

Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 EELTKYGAHVIMLFVPVTLGVVVAIVKSVRFYTERKNGQLIYPTFTDTPSVGQRL 60

|||||

DB 77 EELTKYGAHVIMLFVPVTLGVVVAIVKSVRFYTERKNGQLIYPTFTDTPSVGQRL 136

|||||

QY 61 NSVLTLMISVTVVMTIFLVWLYKYRCYKFIHGWLIMSSLMFLFTYTYLGVGLTKYN 120

|||||

DB 137 NSVLTLMISVTVVMTIFLVWLYKYRCYKFIHGWLIMSSLMFLFTYTYLGVGLTKYN 196

|||||

QY 121 VAMDYPTLLITVYVNFVAVGAVGVCIHWKGLPLVQQAAYLIMISALMALVFIKYLPEWSAVIL 180

|||||

DB 197 VAMDYPTLLITVYVNFVAVGAVGVCIHWKGLPLVQQAAYLIMISALMALVFIKYLPEWSAVIL 256

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QY 181 GAISVYDVLVAVLCPKGLPLMLVETAQERNEPIFPALIYSSAMVYVGMVMAKLDPSQGALQ 240

|||||

Domain 225..244
/label= TM5
/note= "transmembrane domain 5"
245..249
/label= TM5-6
/note= "hydrophilic loop"
250..268
/label= TM6
/note= "transmembrane domain 6"
269..387
/label= TM6-7
/note= "hydrophilic loop"
388..409
/label= TM8
/note= "transmembrane domain 8"
Misc_difference 141
/note= "Asn141Ile mutation site (Claim 19)"
Misc_difference 239
/note= "Met239Val mutation site (Claim 19)"
Misc_difference 420
/note= "Ile420Thr mutation site"
WO9801549-A2.
15-JAN-1998.
04-JUL-1997; CA0475.
02-JAN-1997; US-034590.
05-JUL-1996; US-021673.
12-JUL-1996; US-021700.
08-NOV-1996; US-029895.
(HSCR-) HSC RES & DEV LP.
(UTOR) UNIV TORONTO GOVERNING COUNCIL.
Fraser PE, Rommens JM, St George-Hyslop PH;
WPI: 98-286355/25.
N-PSDB; V04669.
New isolated mutant presenilin-1 genes - useful for developing
products for use in detection, diagnosis and therapy of Alzheimer's
disease and for drug screening
Claim 19: Page 203-204; 238pp; English.
This polypeptide comprises human presenilin-2 (hPS2). Its amino
acid sequence was deduced from an isolated cDNA clone (see V04669).
Human and murine presenilin-1 sequences are also provided (see
W23964-66). Mutations in the PS-1 and PS-2 genes are linked to
the development in humans of forms of familial Alzheimer's disease
(FAD) and may be causative of other disorders, e.g. cognitive,
intellectual, neurological or physiological disorders such as
cerebral haemorrhage, schizophrenia, depression, mental retardation
and epilepsy. Use of the nucleic acids and proteins comprising or
derived from the presenilins is made in screening and diagnosing
FAD, identifying and developing therapeutics for treatment of FAD,
and in producing cell lines and transgenic animals useful as models
of FAD. Methods for identifying substances that bind to, or
modulate the activity of a presenilin protein, and methods for
identifying substances that affect the interaction of a
presenilin-interacting protein with a presenilin protein are also
disclosed.
Sequence 448 AA;

Query Match 100.0%; Score 1923; DB 1; Length 448;
Best Local Similarity 100.0%; Pred. No. 1.6e-200;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 EELTKYGAHVIMLFVPTVLCMIVVATIKSVRFYTEKNGQLIYPTFTDTPSVGQRL 60
Db 77 EELTKYGAHVIMLFVPTVLCMIVVATIKSVRFYTEKNGQLIYPTFTDTPSVGQRL 136
QY 61 NSVLNTLMISVIVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 120
Db 137 NSVLNTLMISVIVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 196
QY 121 VAMDYPTLLLTVMNFGAVGVCVHKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVIL 180
Db 197 VAMDYPTLLLTVMNFGAVGVCVHKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVIL 256

QY 181 GAISVYDLVAVLCPKGPLRMLVETAQERNEPIFPALYSSAMVWTVGMALDPSSQALQ 240
Db 257 GAISVYDLVAVLCPKGPLRMLVETAQERNEPIFPALYSSAMVWTVGMALDPSSQALQ 316
QY 241 LPYDPEMEEDSYDSFGPEPSYPEVPEPLTGYPGEELEEEERGKVLGLGDFIFYSVLVGK 300
Db 317 LPYDPEMEEDSYDSFGPEPSYPEVPEPLTGYPGEELEEEERGKVLGLGDFIFYSVLVGK 376
QY 301 AATGSGDWNNTLACFVAILIGLCLTLLLLAVFKKALPALPISITFTGLIIFYSTDLNVRP 360
Db 377 AATGSGDWNNTLACFVAILIGLCLTLLLLAVFKKALPALPISITFTGLIIFYSTDLNVRP 436
QY 361 FMDTLASHQLYI 372
Db 437 FMDTLASHQLYI 448

RESULT 4
W05763
ID W05763 standard; Protein; 448 AA.
AC W05763;
DT 25-JUL-1997 (first entry)
DE Presenilin-2 M239V mutation.
KW Presenilin-2; human; hPS1-1; hPS1-2; integral membrane protein; AD;
KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;
KW depression; antibody; gene expression modulator; therapy; mutein.
OS Homo sapiens.
FH Key Location/Qualifiers
FT modified_site 239 /label= M239V
FT W09634099-A2.
PN 31-OCT-1996.
PD 29-APR-1996; CA0263.
PR 28-JUN-1995; US-431048.
PR 28-JUN-1995; US-496841.
PR 31-JUL-1995; US-509359.
PA (HSCR-) HSC RES & DEV LP.
PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
Fraser PE, Rommens JM, St George-Hyslop PH;
WPI: 96-497631/49.
PI New presenilin genes - useful for diagnosis, therapy and drug
screening of familial Alzheimer's disease, cerebral disorders, etc.
Claim 4: Page -; 178pp; English.
W05763-W05766 represent mutated versions of the human presenilin-2
protein (see W05762 for wild type sequence). The presenilins are a family
of highly conserved integral membrane proteins with a common structural
motif, common alternate splicing patterns, and common mutational hot spot
regions. Mutations in PS genes are implicated in familial Alzheimer's
disease (AD) and possibly other diseases such as cerebral haemorrhage,
schizophrenia, depression etc., so detection of mutations in the DNA
encoding the wild type sequences can be used for diagnosis of these
diseases. The wild type proteins, or vectors that express them or
containing antisense sequences, antibodies selective for these mutant
forms of the proteins and modulators of PS gene expression are
potentially useful for treatment of AD etc. transgenic animals are useful
as models for drug screening. The antibodies can also be used e.g. for
affinity purification and in immunoassays.
Sequence 448 AA;

Query Match 99.8%; Score 1919; DB 1; Length 448;
Best Local Similarity 99.7%; Pred. No. 4.3e-200;
Matches 371; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 EELTKYGAHVIMLFVPTVLCMIVVATIKSVRFYTEKNGQLIYPTFTDTPSVGQRL 60
Db 77 EELTKYGAHVIMLFVPTVLCMIVVATIKSVRFYTEKNGQLIYPTFTDTPSVGQRL 136
QY 61 NSVLNTLMISVIVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 120
Db 137 NSVLNTLMISVIVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 196
QY 121 VAMDYPTLLLTVMNFGAVGVCVHKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVIL 180

QY 61 NSVLTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSLMFLFTYIYLGEVLKTYN 120
DB 137 NSVLTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSLMFLFTYIYLGEVLKTYN 196
QY 121 VAMDYPTLLTWNFGAVGMYCIHWKGPLVLOQAYLIMISALMALVFTKYLPEWSAWVIL 180
DB 197 VAMDYPTLLTWNFGAVGMYCIHWKGPLVLOQAYLIMISALMALVFTKYLPEWSAWVIL 256
QY 181 GAISVYDLVAVLCPKGPLRMLVETAQRNEPIFFPALIYSSAMWTVGMKLDPSOGALQ 240
DB 257 GAISVYDLVAVLCPKGPLRMLVETAQRNEPIFFPALIYSSAMWTVGMKLDPSOGALQ 316
QY 241 LPYDPEMEEDSYDSFGSPSYPEVFPPLTGYPGEELEEEERGVKLGDFIFYSVLVGK 300
DB 317 LPYDPEMEEDSYDSFGSPSYPEVFPPLTGYPGEELEEEERGVKLGDFIFYSVLVGK 376
QY 301 AAATGSGDNTTACFAVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 360
DB 377 AAATGSGDNTTACFAVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 436
QY 361 FMDTLASHQLYI 372
DB 437 FMDTLASHQLYI 448
RESULT 7
W28508
ID W28508 standard; Protein; 447 AA.
AC W28508;
DE 07-DEC-1997 (first entry)
DT Full AD4/AD3LP sequence.
KW AD3; AD4/AD3LP; Alzheimer's disease; chromosome; missegregation;
KW Presenilin; inhibitor; AD; trisomy 21; ss.
OS Homo sapiens.
PN WO9707213-A2.
PD 27-FEB-1997.
PF 15-AUG-1996; U13314.
PR 16-AUG-1995; US-002448.
PA (HARD) HARVARD COLLEGE.
PI Li J, Potter H;
DR WPI: 97-165297/15.
DR N-PSDB: T87426.
PT Identifying genes which cause chromosome missegregation - useful for
PT identifying causes of and treatments for diseases, e.g. Alzheimer's
PS Claim 29; Fig 29; 77pp; English.
CC Identifying genes which cause improper chromosome segregation,
CC screening for inhibitors of chromosome missegregation and processes
CC caused by genes encoding chromosome missegregation promoters
CC was exemplified using Alzheimer's disease. The sequences
CC given in T87401 to T87426 can be used in the above methods.
CC It is not clear from the figure legend, the figure and the
CC disclosure of the specification which sequence of Fig 1 and Fig 28
CC is the AD4/AD3LP or the AD3 sequence.
SQ Sequence 447 AA;
Query Match 99.2%; Score 1907.5; DB 1; Length 447;
Best Local Similarity 99.7%; Pred. No. 7.6e-199;
Matches 371; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
QY 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFYETKNGQLIYPTPTEDTPSVGQRL 60
DB 77 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFYETKNGQLIYPTPTEDTPSVGQRL 136
QY 61 NSVLTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSLMFLFTYIYLGEVLKTYN 120
DB 137 NSVLTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSLMFLFTYIYLGEVLKTYN 196
QY 121 VAMDYPTLLTWNFGAVGMYCIHWKGPLVLOQAYLIMISALMALVFTKYLPEWSAWVIL 180
DB 197 VAMDYPTLLTWNFGAVGMYCIHWKGPLVLOQAYLIMISALMALVFTKYLPEWSAWVIL 256

QY 181 GAISVYDLVAVLCPKGPLRMLVETAQRNEPIFFPALIYSSAMWTVGMKLDPSOGALQ 240
DB 257 GAISVYDLVAVLCPKGPLRMLVETAQRNEPIFFPALIYSSAMWTVGMKLDPSOGALQ 316
QY 241 LPYDPEMEEDSYDSFGSPSYPEVFPPLTGYPGEELEEEERGVKLGDFIFYSVLVGK 300
DB 317 LPYDPEMEEDSYDSFGSPSYPEVFPPLTGYPGEELEEEERGVKLGDFIFYSVLVGK 375
QY 301 AAATGSGDNTTACFAVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 360
DB 376 AAATGSGDNTTACFAVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 435
QY 361 FMDTLASHQLYI 372
DB 436 FMDTLASHQLYI 447
RESULT 8
W05766
ID W05766 standard; Protein; 414 AA.
AC W05766;
DT 25-JUL-1997 (first entry)
DE Presenilin-2 delta263-296 mutation.
KW Presenilin-2; human; hPS1-1; hPS1-2; PS-2; integral membrane protein; AD;
KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;
KW depression; antibody; gene expression modulator; therapy; mutuin.
OS Homo sapiens.
FH Key
FT Location/Qualifiers
FT misc_difference 263..264
FT /note="site of 34 residue deletion"
PN W09634099-A2.
PD 31-OCT-1996.
PF 29-APR-1996; CA0263.
PR 28-APR-1995; US-431048.
PR 28-JUN-1995; US-496841.
PR 31-JUL-1995; US-509359.
PR (HSCR-) HSC RES & DEV LP.
PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
PI Fraser PE, Rommens JM, St George-Hyslop PH;
DR WPI: 96-497631/49.
DR New presenilin genes - useful for diagnosis, therapy and drug
PT screening of familial Alzheimer's disease, cerebral disorders, etc.
PS Claim 4; Page -; 178pp; English.
CC W05763-W05766 represent mutated versions of the human presenilin-2
CC protein (see W05762 for wild type sequence). The presenilins are a family
CC of highly conserved integral membrane proteins with a common structural
CC motif, common alternate splicing patterns, and common mutational hot spot
CC regions. Mutations in PS genes are implicated in familial Alzheimer's
CC disease (AD) and possibly other diseases such as cerebral haemorrhage,
CC schizophrenia, depression etc., so detection of mutations in the DNA
CC encoding the wild type sequences can be used for diagnosis of these
CC diseases. The wild type proteins, or vectors that express them or
CC containing antisense sequences, antibodies selective for these mutant
CC forms of the proteins and modulators of PS gene expression are
CC potentially useful for treatment of AD etc. Transgenic animals are useful
CC as models for drug screening. The antibodies can also be used e.g. for
CC affinity purification and in immunoassays.
SQ Sequence 414 AA;
Query Match 89.6%; Score 1723; DB 1; Length 414;
Best Local Similarity 90.9%; Pred. No. 7.4e-179;
Matches 338; Conservative 0; Mismatches 0; Indels 34; Gaps 1;
QY 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFYETKNGQLIYPTPTEDTPSVGQRL 60
DB 77 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFYETKNGQLIYPTPTEDTPSVGQRL 136
QY 61 NSVLTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSLMFLFTYIYLGEVLKTYN 120
DB 137 NSVLTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSLMFLFTYIYLGEVLKTYN 196

Qy	241	LPVDPENE-----EDSVDSGEGPSYSEVFPFPPTGTGPG-----EEL-----277
Db	309	VPKNPKYNTQARRETQDSDGNDGGFSEWEAQRDSDLGPHRSTPESRAAVQELSGSI 368
Qy	277	---EEERGVKGLGDFIYSVLVGKAATGSGDNTTILACFVAILIGLCTLLLLAYF 333
Db	369	LTSEDPEERGKVLGDFIYSVLVGKASATAGSDNTTILACFVAILIGLCTLLLLAYF 428
Qy	334	KKALPALPISITFGILFYFSTDMLVRFPMFDTLASHQLYI 372
Db	429	KKALPALPISITFGILFYFATDYLVPFMDQLAFHFYI 467
RESULT	11	
W23966		
ID	W23966	standard; Protein; 467 AA.
AC	W23966;	
DT	20-JUL-1998	(first entry)
DE	Mouse presenilin-1 homologue.	
KW	Presenilin-1; PSI gene; mouse; familial Alzheimer's disease; FAD;	
KW	cerebral haemorrhage; schizophrenia; depression; epilepsy;	
KW	mental retardation; diagnosis; therapy; transgenic animal.	
OS	Mus musculus.	
FH	Key	Location/Qualifiers
FT	Domain	82..100
FT		/label= TM1
FT	Domain	/note= "transmembrane domain 1"
FT		101..132
FT		/label= TM2
FT	Domain	/note= "hydrophilic loop"
FT		133..154
FT		/label= TM2
FT	Domain	/note= "transmembrane domain 2"
FT		155..163
FT		/label= TM2-3
FT	Domain	/note= "hydrophilic loop"
FT		164..183
FT		/label= TM3
FT	Domain	/note= "transmembrane domain 3"
FT		184..194
FT		/label= TM3-4
FT	Domain	/note= "hydrophilic loop"
FT		195..212
FT		/label= TM4
FT	Domain	/note= "transmembrane domain 4"
FT		213..220
FT		/label= TM4-5
FT	Domain	/note= "hydrophilic loop"
FT		221..238
FT		/label= TM5
FT	Domain	/note= "transmembrane domain 5"
FT		239..243
FT		/label= TM5-6
FT	Domain	/note= "hydrophilic loop"
FT		244..262
FT		/label= TM6
FT	Domain	/note= "transmembrane domain 6"
FT		263..407
FT		/label= TM6-7
FT	Domain	/note= "hydrophilic loop"
FT		408..428
FT		/label= TM8
FT	Misc_difference 177	/note= "transmembrane domain 8"
FT		
FT	Misc_difference 439	/note= "Phe177Ser mutation site (Claim 1)"
FT		
FT	Misc_difference 439	/note= "Ile439Val mutation site (Claim 1)"
FT		
PN	WO9801549-A2.	
PN	15-JAN-1998.	
PD	04-JUL-1997.	CA0475.
PR	02-JAN-1997.	US-034590.
PR	05-JUL-1996.	US-021673.
PR	12-JUL-1996.	US-021700.

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08-NOV-1996; US-029895.
PA (HSCR-) HSC RES & DEV LP.
PA (UTOR ) UNIV TORONTO GOVERNING COUNCIL.
PI Fraser PE, Rommens JM, St George-Hyslop PH;
DR WPI; 98-286355/25.
DR N-PSDB; V04668.
PT New isolated mutant presenilin-1 genes - useful for developing
PT products for use in detection, diagnosis and therapy of Alzheimer's
PT disease and for drug screening
PS Disclosure; Page 199-200; 238pp; English.
CC This polypeptide comprises the murine presenilin-1 (PS1) homologue.
CC Its amino acid sequence was deduced from an isolated cDNA clone
CC (see V04668). Mutations in the human PS1 and PS2 genes (see
CC V04665-68) have been linked to the development in humans of forms
CC of familial Alzheimer's disease (FAD). All amino acids that are
CC mutated in analysed FAD pedigrees (see W23964) were conserved in
CC the murine homologue. Use of the nucleic acids and proteins
CC comprising or derived from presenilins can be made in screening and
CC diagnosing FAD, identifying and developing therapeutics for
CC treatment of FAD, and in producing cell lines and transgenic
CC animals useful as models of FAD. Methods for identifying
CC substances that bind to, or modulate the activity of a presenilin
CC protein, and methods for identifying substances that affect the
CC interaction of a presenilin-interacting protein with a presenilin
CC protein are also disclosed.
SQ Sequence 467 AA;

Query Match 74.4%; Score 1431.5; DB 1; Length 467;
Best Local Similarity 71.2%; Pred. No. 4e-147;
Matches 284; Conservative 36; Mismatches 50; Indels 29; Gaps

QY 1 EELFLKYGAHVIMLFPVTLCTMIVVATIKSVRYETKNGQLIYTPFETDPSVGORLL 60
DB 71 EELFLKYGAHVIMLFPVTLCTMIVVATIKSVRYETKNGQLIYTPFETDPSVGORLL 130
QY 61 NSVLTLMISVIVMTIFLWLYKYRCYKFIHGWLIMSMLLFTYIYLGEVLKTYN 120
DB 131 HSLNAAIMISVIVMTILLVLYKYRCYKVIHAWLIISLALLFFSFYILGEVFKTYN 190
QY 121 VAMDYPTLLTVNFGAVGVCIHWKGPLVQQAYLIMISALMALVFIKYLPSWSAWIL 180
DB 191 VADVTVTVALLINFGVVGMIATHWKGPLRQQAYLIMISALMALVFIKYLPEWTAWIL 250
QY 181 GAISYDLDVAVLCPKPLRMVETAQERNPEIPFALLIYSSAWVTYGMALDPSSOGALQ 240
DB 251 AVISYDLDVAVLCPKPLRMVETAQERNPEIPFALLIYSSAWVTYGMALDPSSOGALQ 308
QY 241 LPYDPEME-----EDSYDSFCEPSYVFVFPPLTGVPG-----EEL- 277
DB 309 VPKNPKYNTQARETQDSSGNDGDFSEWAQARDSHLGPHRSTPESRAAVQELS GSI 368
QY 277 ---EEEBEERGKGLGDFIFYSVLVGKAAATGSGDNWNTTIACFVAILIGLCTLLLAIF 333
DB 369 LTSDEPDEERGKGLGDFIFYSVLVGKASATSGDNWNTTIACFVAILIGLCTLLLAIF 428
QY 334 KKALPALPISITFGLIIFYFTSDNMLVRPFMDTSLASHQLYI 372
DB 429 KKALPALPISITFGLVFFATDYLVQPMQDLAFHFQYI 467

RESULT 12
W11839
ID W11839 standard; Protein; 467 AA.
AC W11839;
DT 07-MAY-1997 (first entry)
DE Human early onset Alzheimer's disease (EOAD) polypeptide.
KW Early onset Alzheimer's disease; EOAD; neurodegenerative disease;
KW diagnosis; therapy; inhibitor; antagonist; antibody.
OS Homo sapiens.
FH Key Location/Qualifiers
FT misc_difference 26..29
TT /note= "unidentified amino acid residues"

```

CC alternate splicing of the genomic DNA sequence. W05762 represents the
CC coding sequence for wild type human PS-2. The presenilins are a family of
CC highly conserved integral membrane proteins with a common structural
CC motif, common alternate splicing patterns, and common mutational hot spot
CC regions. Mutations in PS genes are implicated in familial Alzheimer's
CC disease (AD) and possibly other diseases such as cerebral haemorrhage,
CC schizophrenia, depression etc., so detection of mutations in the DNA
CC encoding these sequences can be used for diagnosis of these diseases.
CC These proteins, or vectors that express them or containing antisense
CC sequences, antibodies selective for mutant forms of these proteins (such
CC as W05736) and modulators of PS gene expression are potentially useful
CC for treatment of AD etc. Transgenic animals are useful as models for drug
CC screening. The antibodies can also be used e.g. for affinity purification
CC and in immunoassays.
SQ Sequence 467 AA;

Query Match 74.3%; Score 1429.5; DB 1; Length 467;
Best Local Similarity 72.3%; Pred. No. 6.6e-147;
Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

Qy 1 EELTKYGAHVIMLFVPTLCMVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRL 60
Db 71 EELTKYGAHVIMLFVPTLCMVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRL 130
Qy 61 NSVLTNTLMISVIIVMTIFLVLYKYRCYKFTGHWLMSLMLEFLTYIYLGEVLKTYN 120
Db 131 HSLNAAIMISVIIVMTIFLVLYKYRCYKFTGHWLMSLMLEFLTYIYLGEVLKTYN 190
Qy 121 VANDYPTLLTNVNFAGVGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWTAWL 180
Db 191 VANDYPTLLTNVNFAGVGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWTAWL 250
Qy 181 GAISYDVLVAVLCPCPKPLRMVETAQERNEIFPALLYSSAMVTVGMKLDPSOGAL- 240
Db 251 AVISYDVLVAVLCPCPKPLRMVETAQERNEIFPALLYSSAMVTVGMKLDPSOGAL- 310
Qy 240 -QLPYDPE-MEEDSYDSFGE---PSYPVEFPPLTGYPG-----EEL----- 277
Db 311 KNSKYNASTERESQDTVAENDDGGFSEWEAQRDHLGPHRSTPSRAAVQELSSILA 370
Qy 277 -EEEEERGKVLGLGDFIFYSVLVGKAAATGSGDWNNTTACFVAILIGLCLTLLLAIFKK 335
Db 371 GEDPEERGKVLGLGDFIFYSVLVGKASATASGDWNNTTACFVAILIGLCLTLLLAIFKK 430
Qy 336 ALPALPISITFGLIFYFSTDNLVRPMDTLASHQLYI 372
Db 431 ALPALPISITFGLIFYFATDYLVPQMDQLAFHQFYI 467

RESULT 15
W28507
ID W28507 standard; Protein; 407 AA.
AC W28507;
DT 07-DEC-1997 (first entry)
DE Partial AD3 sequence.
KW AD3; AD4/AD3LP; Alzheimer's disease; chromosome; missegregation;
KW presenilin; inhibitor; AD; trisomy 21.
OS Homo sapiens.

FT Key Location/Qualifiers
FT misc_difference 86
FT /label= mutation
FT /note= "M -> L"
FT misc_difference 103
FT /label= mutation
FT /note= "H -> R"
FT misc_difference 186
FT /label= mutation
FT /note= "A -> E"
FT misc_difference 226
FT /label= mutation
FT /note= "L -> V"

FT /label= mutation
FT /note= "C -> Y"
PN W09707213-A2.
PD 27-FEB-1997.
PF 15-AUG-1996; U13314.
PR 16-AUG-1995; US-002448.
PA (HARD) HARVARD COLLEGE.
PI Li J, Potter H;
DR WPI; 97-165297/15.
DR N-PSDB; T87402.
PT Identifying genes which cause chromosome missegregation - useful for
PT identifying causes of and treatments for diseases, e.g. Alzheimer's
PT disease, cancer and ageing
PS Disclosure; Fig 1; 77pp; English.
CC Identifying genes which cause improper chromosome segregation,
CC screening for inhibitors of chromosome missegregation and processes
CC caused by genes encoding chromosome missegregation promoters
CC was exemplified using Alzheimer's disease. The sequences
CC given in T87401 to T87426 can be used in the above methods.
CC The five mutations indicated in the Features Table cosegregate
CC with early-onset familial Alzheimer's disease. It is predicted
CC that these mutations result in increased levels of cells with
CC trisomy 21 in carriers of the mutation compared with non-carriers.
SQ Sequence 407 AA;

Query Match 74.3%; Score 1429.5; DB 1; Length 407;
Best Local Similarity 72.3%; Pred. No. 5.4e-147;
Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

Qy 1 EELTKYGAHVIMLFVPTLCMVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRL 60
Db 11 EELTKYGAHVIMLFVPTLCMVVATIKSVRYTEKNGQLIYPTPTDTPSVGQRL 70
Qy 61 NSVLTNTLMISVIIVMTIFLVLYKYRCYKFTGHWLMSLMLEFLTYIYLGEVLKTYN 120
Db 71 HSLNAAIMISVIIVMTIFLVLYKYRCYKFTGHWLMSLMLEFLTYIYLGEVLKTYN 130
Qy 121 VANDYPTLLTNVNFAGVGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWTAWL 180
Db 131 VANDYPTLLTNVNFAGVGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWTAWL 190
Qy 181 GAISYDVLVAVLCPCPKPLRMVETAQERNEIFPALLYSSAMVTVGMKLDPSOGAL- 240
Db 191 AVISYDVLVAVLCPCPKPLRMVETAQERNEIFPALLYSSAMVTVGMKLDPSOGAL- 250
Qy 240 -QLPYDPE-MEEDSYDSFGE---PSYPVEFPPLTGYPG-----EEL----- 277
Db 251 KNSKYNASTERESQDTVAENDDGGFSEWEAQRDHLGPHRSTPSRAAVQELSSILA 310
Qy 277 -EEEEERGKVLGLGDFIFYSVLVGKAAATGSGDWNNTTACFVAILIGLCLTLLLAIFKK 335
Db 311 GEDPEERGKVLGLGDFIFYSVLVGKASATASGDWNNTTACFVAILIGLCLTLLLAIFKK 370
Qy 336 ALPALPISITFGLIFYFSTDNLVRPMDTLASHQLYI 372
Db 371 ALPALPISITFGLIFYFATDYLVPQMDQLAFHQFYI 407

Search completed: March 20, 2000, 05:31:21
Job time: 4210 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 19:55:31 ; Search time 26.47 seconds
(without alignments)
186.879 Million cell updates/sec

Title: US-08-509-359B-138

Perfect score: 1923

Sequence: 1 EELTKYGAHVIMLFVPT.....STDNLVRFPMDTLASHQLYI 372

Scoring table: BLOSUM62

Searched: 133990 seqs, 13297546 residues

Database : Issued_Patents_AA.*

Word size : 0

Number of hits that pass the threshold : 133990
1: /cgn2_6/ptodata/2/iaa/5A_COMB.pep.*
2: /cgn2_6/ptodata/2/iaa/5B_COMB.pep.*
3: /cgn2_6/ptodata/2/iaa/6_COMB.pep.*
4: /cgn2_6/ptodata/2/iaa/PCTUS9_COMB.pep.*
5: /cgn2_6/ptodata/2/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1923	100.0	448	2	US-08-967-101-137
2	1923	100.0	372	2	US-08-967-101-138
3	1923	100.0	448	2	US-08-592-541-137
4	1923	100.0	372	2	US-08-592-541-138
5	1907.5	99.2	447	2	US-08-875-972-29
6	1591.5	82.8	376	2	US-08-875-972-2
7	1429.5	74.3	407	2	US-08-967-101-134
8	1429.5	74.3	407	2	US-08-875-972-4
9	1429.5	74.3	467	2	US-08-592-541-134
10	1429.5	74.3	467	3	US-08-670-964-2
11	1429.5	74.3	463	3	US-08-670-964-4
12	1423.5	74.0	467	2	US-08-967-101-2
13	1423.5	74.0	467	2	US-08-592-541-2
14	1412.5	73.5	463	2	US-08-670-479-18
15	1381.5	71.8	467	2	US-08-967-101-4
16	1381.5	71.8	467	2	US-08-592-541-4
17	1127.5	58.6	541	2	US-08-967-101-166
18	1127.5	58.6	541	2	US-08-592-541-166
19	100	5.2	1294	2	US-08-819-288-3
20	98	5.1	1321	1	US-08-261-822A-3
21	98	5.1	1321	4	PCT-US95-0774A-3
22	97	5.0	1334	2	US-08-996-345-2
23	89	4.6	509	2	US-09-031-392-6
24	88.5	4.6	452	1	US-08-117-361C-1
25	88.5	4.6	3169	2	US-08-477-451-6
26	87	4.5	492	2	US-08-355-844-3
27	87	4.5	492	4	PCT-US95-16126-3
28	82	4.3	502	1	US-08-484-840-3
29	82	4.3	1873	1	US-08-336-257A-7
30	82	4.3	502	1	US-08-483-094-3
31	82	4.3	413	2	US-08-808-793-25
32	81	4.2	2100	2	US-08-808-793-23
33	80	4.2	837	1	US-07-923-976-2
34	80	4.2	602	1	US-08-295-814E-2

ALIGNMENTS

RESULT 1

US-08-967-101-137
; Sequence 137, Application US/08967101
; Patent No. 5840340

; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; FILING DATE: 10-NOV-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/592,541
; FILING DATE:

; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100

; INFORMATION FOR SEQ ID NO: 137:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 448 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear

; MOLECULE TYPE: protein
US-08-967-101-137

Query Match 100.0%; Score 1923; DB 2; Length 448;
Best Local Similarity 100.0%; Pred. No. 2.2e-190;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFTEKNGQLIYPTFTDTPSVGQRL 60

Db 77 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFTEKNGQLIYPTFTDTPSVGQRL 136

Qy 61 NSVLATLIMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 120

Db 137 NSVLATLIMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLMFLFTYIYLGEVLKTYN 196

Db 77 EELTKYGAHVIMLFVPTLCMIVVWATIKSVRFYTEKNGQLIYTPFTEDPSVGQRL 136
QY 61 NSVLTLMISVIVVMTFLVLYKYCYKFIHGWLIMSSLMFLFYIYLGEVLKTYN 120
Db 137 NSVLTLMISVIVVMTFLVLYKYCYKFIHGWLIMSSLMFLFYIYLGEVLKTYN 196
QY 121 VMDYPTLLTWNFGAVGWCIVHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWIL 180
Db 197 VMDYPTLLTWNFGAVGWCIVHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWIL 256
QY 181 GAISYDYLAVLCPKGPLRMLVETAQERNEPIFPALIIYSSAMVTVGMKLDPSQGALQ 240
Db 257 GAISYDYLAVLCPKGPLRMLVETAQERNEPIFPALIIYSSAMVTVGMKLDPSQGALQ 316
QY 241 LPYDPEMEDSDYSGEPSPYEVFEPPLTGYGPEEEERGVKLGDFIFYSVLVGK 300
Db 317 LPYDPEMEDSDYSGEPSPYEVFEPPLTGYGPEEEERGVKLGDFIFYSVLVGK 376
QY 301 AAATGSGDWNNTLACFVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 360
Db 377 AAATGSGDWNNTLACFVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 436
QY 361 FMDTLASHOLYI 372
Db 437 FMDTLASHOLYI 448

RESULT 4
US-08-592-541-138
; Sequence 138, Application US/08592541
; Patent No. 5986054
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/592,541
; FILING DATE:
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 138:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 372 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-592-541-138

Query Match 100.0%; Score 1923; DB 2; Length 372;
Best Local Similarity 100.0%; Pred. No. 1.7e-190;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 EELTKYGAHVIMLFVPTLCMIVVWATIKSVRFYTEKNGQLIYTPFTEDPSVGQRL 60
Db 1 EELTKYGAHVIMLFVPTLCMIVVWATIKSVRFYTEKNGQLIYTPFTEDPSVGQRL 60
QY 61 NSVLTLMISVIVVMTFLVLYKYCYKFIHGWLIMSSLMFLFYIYLGEVLKTYN 120
Db 61 NSVLTLMISVIVVMTFLVLYKYCYKFIHGWLIMSSLMFLFYIYLGEVLKTYN 120
QY 121 VMDYPTLLTWNFGAVGWCIVHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWIL 180
Db 121 VMDYPTLLTWNFGAVGWCIVHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWIL 180
QY 181 GAISYDYLAVLCPKGPLRMLVETAQERNEPIFPALIIYSSAMVTVGMKLDPSQGALQ 240
Db 181 GAISYDYLAVLCPKGPLRMLVETAQERNEPIFPALIIYSSAMVTVGMKLDPSQGALQ 240
QY 241 LPYDPEMEDSDYSGEPSPYEVFEPPLTGYGPEEEERGVKLGDFIFYSVLVGK 300
Db 241 LPYDPEMEDSDYSGEPSPYEVFEPPLTGYGPEEEERGVKLGDFIFYSVLVGK 300
QY 301 AAATGSGDWNNTLACFVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 360
Db 301 AAATGSGDWNNTLACFVAILIGLCTLLLLAVFKKALPALPISITFGLIFYFSTDNLVRP 360
QY 361 FMDTLASHOLYI 372
Db 361 FMDTLASHOLYI 372

RESULT 5
US-08-875-972-29
; Sequence 29, Application US/08875972
; Patent No. 5985564
; GENERAL INFORMATION:
; APPLICANT: Huntington Potter and Jinhue Li
; TITLE OF INVENTION: ASSAY FOR IDENTIFYING GENES CAUSING
; TITLE OF INVENTION: CHROMOSOME NON-DISJUNCTION
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HAMILTON, BROOK, SMITH & REYNOLDS, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02173-4799
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/875,972
; FILING DATE: 08-AUG-97
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/002,448
; FILING DATE: 16-AUG-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan Esq., Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: HU95-03PA
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781) 861-6240
; TELEFAX: (781) 861-9540
; INFORMATION FOR SEQ ID NO: 29:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 447 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-875-972-29

; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-967-101-134

Query Match 74.3%; Score 1429.5; DB 2; Length 467;
Best Local Similarity 72.3%; Pred. No. 2e-139;
Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

QY 1 BELTLKYGAKHVMFLFVPVTLCMVVVATIKSVRYTEKNGQLIYTPPTDTPSVGQRL 60
DB 71 BELTLKYGAKHVMFLFVPVTLCMVVVATIKSVRYTEKNGQLIYTPPTDTPSVGQRL 130
QY 61 NSVLNTLMISIVVMTIFLVLYKYRCYKFIHGWLMSSLMFLFYIYLGEVLKTYN 120
DB 131 HSLNAAIMISIVVMTILLVLYKYRCYKVIHAWLIISLSSLLFFSYIYLGEVFKTYN 190
QY 121 VAMDYPTLLTWNFGAVGMCVHMKGPLVLOQAYLIMISALMALVFYKYLPEWSAWIL 180
DB 191 VADYITVALLIWNFGVGMISIHMKGPLRLOQAYLIMISALMALVFYKYLPEWTAWIL 250
QY 181 GAISYDLVAVLCPKGPLRMLVETAQERNETFPALIISSANVTVGMKLDPSOGAL- 240
DB 251 AVISYDLVAVLCPKGPLRMLVETAQERNETFPALIISSANVTVGMKLDPSOGAL- 310
QY 240 -QLPYDPE-MEEDSYDSFGE---PSYPEVFPPLTGYPG-----EEL----- 277
DB 311 KNSKYNAESTERESQDTVAENDDGGFSEWEAQRDHSHLPHRSTPESRAAVQELSSILA 370
QY 277 -EEERGVKGLGDFIFYSVLVKGAAATSGDNTTACFVAILLIGLCLTLLLAIFEKK 335
DB 371 GEDPEERGKVLGDFIFYSVLVKGASATASGDNTTACFVAILLIGLCLTLLLAIFEKK 430
QY 336 ALPALPISITFGLIFYSTDLNLRPFMDTLASHOLYI 372
DB 431 ALPALPISITFGLIFYATDYLVPQFMDQLAFHFYI 467

RESULT 8

US-08-875-972-4
; Sequence 4, Application US/08875972
; Patent No. 5985564

; GENERAL INFORMATION:
; APPLICANT: Huntington Potter and Jinhue Li
; TITLE OF INVENTION: ASSAY FOR IDENTIFYING GENES CAUSING
; TITLE OF INVENTION: CHROMOSOME NON-DISJUNCTION
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HAMILTON, BROOK, SMITH & REYNOLDS, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02173-4799

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; APPLICATION NUMBER: US/08/875,972
; FILING DATE: 08-AUG-97
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/002,448
; FILING DATE: 16-AUG-1995

; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan Esq., Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: HU95-03PA
; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (781) 861-6240
; TELEFAX: (781) 861-9540
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 407 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-875-972-4

Query Match 74.3%; Score 1429.5; DB 2; Length 407;
Best Local Similarity 72.3%; Pred. No. 1.6e-139;
Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

QY 1 BELTLKYGAKHVMFLFVPVTLCMVVVATIKSVRYTEKNGQLIYTPPTDTPSVGQRL 60
DB 11 BELTLKYGAKHVMFLFVPVTLCMVVVATIKSVRYTEKNGQLIYTPPTDTPSVGQRL 70
QY 61 NSVLNTLMISIVVMTIFLVLYKYRCYKFIHGWLMSSLMFLFYIYLGEVLKTYN 120
DB 71 HSLNAAIMISIVVMTILLVLYKYRCYKVIHAWLIISLSSLLFFSYIYLGEVFKTYN 130
QY 121 VAMDYPTLLTWNFGAVGMCVHMKGPLVLOQAYLIMISALMALVFYKYLPEWSAWIL 180
DB 131 VADYITVALLIWNFGVGMISIHMKGPLRLOQAYLIMISALMALVFYKYLPEWTAWIL 190
QY 181 GAISYDLVAVLCPKGPLRMLVETAQERNETFPALIISSANVTVGMKLDPSOGAL- 240
DB 191 AVISYDLVAVLCPKGPLRMLVETAQERNETFPALIISSANVTVGMKLDPSOGAL- 250
QY 240 -QLPYDPE-MEEDSYDSFGE---PSYPEVFPPLTGYPG-----EEL----- 277
DB 251 KNSKYNAESTERESQDTVAENDDGGFSEWEAQRDHSHLPHRSTPESRAAVQELSSILA 310
QY 277 -EEERGVKGLGDFIFYSVLVKGAAATSGDNTTACFVAILLIGLCLTLLLAIFEKK 335
DB 311 GEDPEERGKVLGDFIFYSVLVKGASATASGDNTTACFVAILLIGLCLTLLLAIFEKK 370
QY 336 ALPALPISITFGLIFYSTDLNLRPFMDTLASHOLYI 372
DB 371 ALPALPISITFGLIFYATDYLVPQFMDQLAFHFYI 407

RESULT 9

US-08-592-541-134
; Sequence 134, Application US/08592541
; Patent No. 5986054

; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/592,541
; FILING DATE:
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.

; OPERATING SYSTEM: DOS
 ; SOFTWARE: FastSeq for Windows Version 2.0
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/670,964
 ; FILING DATE: 26-JUN-1996
 ; CLASSIFICATION: 435
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: 60/001,142
 ; FILING DATE: 13-JUL-1995
 ; APPLICATION NUMBER: 60/001,501
 ; FILING DATE: 18-JUL-1995
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Han, William T
 ; REGISTRATION NUMBER: 34,344
 ; REFERENCE/DOCKET NUMBER: P50358
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: 610-270-5219
 ; TELEFAX: 610-270-5090
 ; TELEX:
 ; INFORMATION FOR SEQ ID NO: 4:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 463 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: protein
 ; US-08-670-964-4

Query Match 74.3%; Score 1429.5; DB 3; Length 463;
 Best Local Similarity 72.3%; Pred. No. 1.9e-139;
 Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

QY 1 BELTLKYGAKHVMFLFVPVTLGMVVVATIKSVRYTERKNGOLIYTPFTEDTPSVGQRL 60
 DB 67 BELTLKYGAKHVMFLFVPVTLGMVVVATIKSVRYTERKNGOLIYTPFTEDTPSVGQRL 126
 QY 61 NSVLTLMISIVVMTFLVLYKYRCYKFIHGLWLSMLLFLFYIYLGVEVKYN 120
 DB 127 HSILNAAMISIVVMTILLVLYKYRCYKFIHAWLIISLLELFFSFYILGEVFKYN 186
 QY 121 VAMDYPTLLTWNFGAVGMVCIHMKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVIL 180
 DB 187 VAVDYITVALLIWNFGVGMISIHMKGPLRLOQAYLIMISALMALVFIKYLPEWTAWLIL 246
 QY 181 GAISYDVLVAVLCPRGLRMLVETAQERNEPIFPALIISSAMVWTVGMKLDPSQOGAL- 240
 DB 247 AVISYDVLVAVLCPRGLRMLVETAQERNEPIFPALIISSAMVWTVGMKLDPSQOGAL- 306
 QY 240 -QLPDPE-MEEDSDSFGE---PSYPEVFEPPLTGYPG-----BEL----- 277
 DB 307 KNSKYNAESTERESODTVAENDDGGFSEWEAQRDHLGPHRSTPESRAAVQELSSILA 366
 QY 277 -EEEREGVKLGDFIFYSVLVGKAAATGSGDWNITLACFVAILIGLCTLLLLAVFKK 335
 DB 367 GEDPEERGKLGDFIFYSVLVGKASATASGDWNITLACFVAILIGLCTLLLLAIFKK 426
 QY 336 ALPALPISITFGLIFFSTDLNLRVPMFTLASHOLYI 372
 DB 427 ALPALPISITFGLVFYFATDYLQVFPMDQLAFHQFYI 463

RESULT 12
 US-08-967-101-2
 ; Sequence 2, Application US/08967101
 ; Patent No. 5840540
 ; GENERAL INFORMATION:
 ; APPLICANT: ST. GEORGE-HYSLOP, PETER H
 ; APPLICANT: ROMMENS, JOHANNA M
 ; APPLICANT: FRASER, PAUL E
 ; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 ; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
 ; NUMBER OF SEQUENCES: 183

; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
 ; STREET: High Street Tower - 125 High Street
 ; CITY: Boston
 ; STATE: Massachusetts
 ; COUNTRY: U.S.A.
 ; ZIP: 02110
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: PatentIn Release #1.0, Version #1.30
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/967,101
 ; FILING DATE: 10-NOV-1997
 ; CLASSIFICATION: 435
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: 08/592,541
 ; FILING DATE:
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Fitcher, Edmund R.
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (617) 248-7000
 ; TELEFAX: (617) 248-7100
 ; INFORMATION FOR SEQ ID NO: 2:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 467 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: protein
 ; US-08-967-101-2

Query Match 74.0%; Score 1423.5; DB 2; Length 467;
 Best Local Similarity 72.0%; Pred. No. 8.2e-139;
 Matches 286; Conservative 33; Mismatches 53; Indels 25; Gaps 5;

QY 1 BELTLKYGAKHVMFLFVPVTLGMVVVATIKSVRYTERKNGOLIYTPFTEDTPSVGQRL 60
 DB 71 BELTLKYGAKHVMFLFVPVTLGMVVVATIKSVRYTERKNGOLIYTPFTEDTPSVGQRL 130
 QY 61 NSVLTLMISIVVMTFLVLYKYRCYKFIHGLWLSMLLFLFYIYLGVEVKYN 120
 DB 131 HSILNAAMISIVVMTILLVLYKYRCYKFIHAWLIISLLELFFSFYILGEVFKYN 190
 QY 121 VAMDYPTLLTWNFGAVGMVCIHMKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVIL 180
 DB 191 VAVDYITVALLIWNFGVGMISIHMKGPLRLOQAYLIMISALMALVFIKYLPEWTAWLIL 250
 QY 181 GAISYDVLVAVLCPRGLRMLVETAQERNEPIFPALIISSAMVWTVGMKLDPSQOGAL- 240
 DB 251 AVISYDVLVAVLCPRGLRMLVETAQERNEPIFPALIISSAMVWTVGMKLDPSQOGAL- 310
 QY 240 -QLPDPE-MEEDSDSFGE---PSYPEVFEPPLTGYPG-----BEL----- 277
 DB 311 KNSKYNAESTERESODTVAENDDGGFSEWEAQRDHLGPHRSTPESRAAVQELSSILA 370
 QY 277 -EEEREGVKLGDFIFYSVLVGKAAATGSGDWNITLACFVAILIGLCTLLLLAVFKK 335
 DB 371 GEDPEERGKLGDFIFYSVLVGKASATASGDWNITLACFVAILIGLCTLLLLAIFKK 430
 QY 336 ALPALPISITFGLIFFSTDLNLRVPMFTLASHOLYI 372
 DB 431 ALPALPISITFGLVFYFATDYLQVFPMDQLAFHQFYI 467

RESULT 13
 US-08-592-541-2
 ; Sequence 2, Application US/08592541
 ; Patent No. 5986054
 ; GENERAL INFORMATION:
 ; APPLICANT: ST. GEORGE-HYSLOP, PETER H

Db 367 GEDPEERGKGLGDFIFYSVLGKASATASGDWNTTIACFVAILIGLCLTLLLAIFKK 426
Qy 336 ALPALPISITGLFIFSTDNIVRFMDTLASHQIYI 372
Db 427 ALPALPISITGLFVFYFATDYLQVPMQDLAFHQFYI 463
RESULT 15
US-08-967-101-4
; Sequence 4, Application US/08967101
; Patent No. 5840540
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/967,101
; FILING DATE: 10-NOV-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/592,541
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 467 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-967-101-4

Query Match 71.8%; Score 1381.5; DB 2; Length 467;
Best Local Similarity 68.7%; Pred. No. 1.8e-134;
Matches 274; Conservative 38; Mismatches 58; Indels 29; Gaps 4;
Qy 1 BELTLKYGAKHIVLFPVTLICMIVVATIKSVRYTEKNGOLIYTPETDTPSYGQRL 60
Db 71 BELTLKYGAKHIVLFPVTLICMIVVATIKSVRYTEKNGOLIYTPETDTPSYGQRL 130
Qy 61 NSVLNTLMISVIVMTFLVLYKYRCYKFIHGLWLMSSMLLFLFYIYLGEVLYKTYN 120
Db 131 HSILNAAIMISVIVMTFLVLYKYRCYKFIHGLWLMSSMLLFLFYIYLGEVLYKTYN 190
Qy 121 VMDYPTLLLTWNFGAVCMCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWIL 180
Db 191 VXDVTVTALLIWNWVGVIHAIHWKGPLRLOQAYLIMISALMALVFIKYLPEWTAWIL 250
Qy 181 GAISYVDLVAVLCPKGPLRMVETAEQNEPIFFPALIYSSAMVTVGMKLDPPSSOGALQ 240
Db 251 AVISYVDLVAVLCPKGPLRMVETAEQNEPIFFPALIYSSAMVTVGMKLDPPSSOGALQ 308

Qy 241 LPYDPEME-----EDSYDSFGSPSYEVEPEPLITGYPG-----EEL----- 277
Db 309 YPKNPKYNTQRAERTQDSGSGNDGSGFSEWEAQDSDHLGPHRSTPESRAAVQELSGSI 368
Qy 277 ---EEEEERGVKGLGDFIFYSVLGKAAATGSGDWNTTIACFVAILIGLCLTLLLAIF 333
Db 369 LTSDEPPEERGVKGLGDFIFYSVLGKASATASGDWNTTIACXVAILIGLCLLALLAIY 428
Qy 334 KKALPALPISITGLFIFSTDNIVRPFMDTLASHQIYI 372
Db 429 KKGXPAXPISITGLFVFYFATDYLQVPMQDLAFHQFYI 467

Search completed: March 18, 2000, 19:55:32
Job time: 3230 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 14:11:52 ; Search time 41.25 Seconds
(without alignments)
425.381 Million cell updates/sec

Title: US-08-509-359B-138
Perfect score: 1923
Sequence: 1 EELTLKYGAKHVIMLFVPT.....STDNLVRPFMDTLASHQLYI 372

Scoring table: BLOSUM62

Searched: 142080 seqs, 47169319 residues

Database : PIR_62.*

Word size : 0

Number of hits that pass the threshold : 142080

1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	1923	100.0	448	2	I58098	E5-1 protein - hum
2	1915	99.6	448	2	A56993	presenilin 2 - hum
3	1802.5	93.7	442	2	J39174	seven trans-membra
4	1534	79.8	449	2	J5391	presenilin-beta -
5	1431.5	74.4	467	2	I78388	S182 protein - mou
6	1431.5	74.4	433	2	JC5390	presenilin-alpha -
7	1429.5	74.3	467	2	S58396	presenilin 1, spli
8	1429.5	74.3	463	2	S63683	presenilin I-463 -
9	1413.5	73.5	463	2	JC5081	presenilin 1 prote
10	1413.5	73.5	467	2	JC5080	presenilin 1 prote
11	1011	52.6	374	2	S63684	presenilin 1, spli
12	956	49.7	461	2	S60253	sel-12 protein - C
13	524.5	27.3	358	2	T15184	hypothetical prote
14	513	26.7	453	2	T00724	presenilin homolog
15	274	14.2	465	2	A43459	sperm membrane pro
16	110	5.7	2016	2	A38195	sodium channel pro
17	104	5.4	826	2	T02268	potassium transpor
18	104	5.4	398	2	H75043	mg2+ transport pro
19	103	5.4	382	2	S47882	ubiquinol--cytochr
20	102	5.3	379	2	I48135	ubiquinol--cytochr
21	101.5	5.3	1681	2	A55138	sodium channel mna
22	100.5	5.2	324	2	S36646	integrin-associate
23	99.5	5.2	1840	1	CHRMI	sodium channel pro
24	99.5	5.2	379	2	JC6178	serotonin receptor
25	99	5.1	2019	2	A33996	sodium channel pro
26	98.5	5.1	531	2	T11596	hypothetical prote
27	98	5.1	461	2	T11829	NADH dehydrogenase
28	98	5.1	217	2	S01095	hypothetical prote
29	97	5.0	381	2	T11440	ubiquinol--cytochr
30	97	5.0	447	2	S52968	NADH dehydrogenase
31	97	5.0	238	2	S02063	H+-transporting AT
32	97	5.0	441	2	S13425	endothelin recepto
33	97	5.0	299	2	D65187	hypothetical 33.7
34	96.5	5.0	308	2	S22328	ubiquinol--cytochr
35	96.5	5.0	1070	2	S25834	rad3 protein - fis

36 96 5.0 507 2 B64433 probable O-antigen
37 96 5.0 592 2 E70488 cytochrome-c oxida
38 95.5 5.0 488 1 QXASM4 NADH dehydrogenase
39 95.5 5.0 442 1 JQ1042 endothelin recepto
40 95.5 5.0 294 2 D53290 oligopeptide trans
41 95.5 5.0 420 2 A47649 probable inner mem
42 95.5 5.0 768 2 S52684 probable membrane
43 95.5 5.0 799 2 H71255 probable cell divi
44 95 4.9 492 2 A30797 glucose transport
45 95 4.9 484 2 S75022 bacteriochlorophyl

ALIGNMENTS

RESULT 1
I58098
E5-1 protein - human
C:Species: Homo sapiens (man)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 29-Sep-1999
C:Accession: I58098
R:Rogaev, E.I.; Sherrington, R.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; Liang, Y.; C
.; Cohen, D.; Lannfelt, L.; Fraser, P.E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 376, 775-778, 1995
A:Title: Familial Alzheimer's disease in kindreds with missense mutations in a gene o
A:Reference number: I58098; MUID:95379971
A:Accession: I58098
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-448 <RES>
A:Cross-references: GB:L44577; NID:g950347; PIDN:AAC42012.1; PID:g950348
C:Genetics:
A:Gene: E5-1
C:Superfamily: presenilin

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Best Local Similarity 100.0%; Pred. No. 3.5e-138;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 EELTLKYGAKHVIMLFVPTLCMIVVATIKSVRYTEKNGOLIYTPFTEDTPSVGQRL 60
DB 77 EELTLKYGAKHVIMLFVPTLCMIVVATIKSVRYTEKNGOLIYTPFTEDTPSVGQRL 136
QY 61 NSVLTLMISVIVVMTIFLVVLYKYRCYKFTHGWLINSSMLLFLFYIYLGEVLKTYN 120
DB 137 NSVLTLMISVIVVMTIFLVVLYKYRCYKFTHGWLINSSMLLFLFYIYLGEVLKTYN 196
QY 121 VMDYPTLLLTWNFGVGMVCIHWKGPLVLOQAAYLIMISALMALVFIKYLPEWSAWVIL 180
DB 197 VMDYPTLLLTWNFGVGMVCIHWKGPLVLOQAAYLIMISALMALVFIKYLPEWSAWVIL 256
QY 181 GAISYDYLAVLCPKPLRMVETAQERNEFPFALIVSSAMVTVGMKLPDSSOGALQ 240
DB 257 GAISYDYLAVLCPKPLRMVETAQERNEFPFALIVSSAMVTVGMKLPDSSOGALQ 316
QY 241 LPYPMEEDSYDSFGESYPEVFPPLTGTYPGELEEEERGVKLGDFIFYSVLVGK 300
DB 317 LPYPMEEDSYDSFGESYPEVFPPLTGTYPGELEEEERGVKLGDFIFYSVLVGK 376
QY 301 AAATGSGDWTTLACFAVAILGLCTLALLAVFKKALPALPISITFGLIFYFSTDNLVRP 360
DB 377 AAATGSGDWTTLACFAVAILGLCTLALLAVFKKALPALPISITFGLIFYFSTDNLVRP 436
QY 361 FMDTLASHQLYI 372
DB 437 FMDTLASHQLYI 448

RESULT 2
A56993
presenilin 2 - human
N:Alternate names: Alzheimer's disease protein 4

QY 240 -QLPY-----DPEMEDSYDSFGPEPSYVEFPPLTGYGPELEEEERGVKLGIDRFY 294
Db 319 QOVQHIDNTPEGANSIVDEAAETRIQ-----TQSNLSEDDPEERGVKLGIDRFY 371
QY 295 SVLVGKAAATGSGDWNNTLACFAVAILGICLTLTLLAVFKKALPALPISITIFGLIFYST 354
Db 372 SVLVGKAAATGSGDWNNTLACFAVAILGICLTLTLLAVFKKALPALPISITIFGLIFYST 431
QY 355 DNLVRPFMDTLASHQLYI 372
Db 432 DNLVRPFMDTLASHQMYI 449
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I78388
S182 protein - mouse
C:Species: Mus musculus (house mouse)
C:Date: 27-Feb-1997 #sequence_revision 27-Feb-1997 #text_change 29-Sep-1999
C:Accession: I78388
R:Sherrington, R.; Rogaev, E.I.; Liang, Y.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; Chl.
ero, I.; Pinessi, L.; Nee, L.; Chumakov, I.; Pollen, D.; Brookes, A.; Sanseau, P.; Polin
E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 375, 754-760, 1995
A:Title: Cloning of a gene bearing missense mutations in early-onset familial Alzheimer
A:Reference number: I58095; MUID:95319502
A:Accession: I78388
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-467 <RES>
A:Cross-references: GB:L42177; NID:g904129; PIDN:AAC42094.1; PID:g904130
C:Superfamily: presenilin
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Best Local Similarity 71.2%; Pred. No. 5.1e-101;
Matches 284; Conservative 36; Mismatches 50; Indels 29; Gaps 4;
QY 1 BELTLKYGAKHVMFLFVPVTLICMIVVATIKSVRYTEKNGQLIYTPPTEDTPSVGQRL 60
Db 71 BELTLKYGAKHVMFLFVPVTLICMIVVATIKSVSFYTRKDGQLIYTPPTEDTPVGQRL 130
QY 61 NSVLNTLMISVIVVMTIFLVLYKYCYKFIHGWLMISLMFLFYIYLGVEVLTYN 120
Db 131 HSLNAAIMSVIVMTILLVLYKYCYKFIHAWLISSLLLFYFIFGLVEVFTYN 190
QY 121 VAMDYPTLLTVMNFGAVGVCIHKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVIL 180
Db 191 VADYVTVALLIWNFGVGMIAIHWKGPLRLOQAYLIMISALMALVFIKYLPEWTAWLIL 250
QY 181 GAISYDVLAVLCPKGLRMLVETAQERNEPIFPALIIYSAMVTVGMKLDPSQSGALQ 240
Db 251 AVISYDVLAVLCPKGLRMLVETAQERNETLFPALIIYSSTMWLVNMAEGDPEAQ--RR 308
QY 241 LPYDPENE-----EDSDSFCEPSYVEFPPLGYPG-----BEL----- 277
Db 309 VPKNPKYNTQARERTQDSGNGDDGFSSEWQAQRDHLGPHRSTPESRAAVQELSGSI 368
QY 277 ---EEERGVKLGIDFIFYSVLVGKAAATGSGDWNNTLACFAVAILGICLTLTLLAVF 333
Db 369 LTSEDPERGVKLGIDFIFYSVLVGKASATASGDWNTTACFAVAILGICLTLTLLAIF 428
QY 334 KKALPALPISITIFGLIFYSTDNLRPFMDTLASHQLYI 372
Db 429 KKALPALPISITIFGLVFFATDYLVPFMDQLAFHQFYI 467
RESULT 6
JC5390
presenilin-alpha - African clawed frog
C:Species: Xenopus laevis (African clawed frog)
C:Date: 04-Jun-1997 #sequence_revision 18-Jul-1997 #text_change 29-Sep-1999
C:Accession: JC5390
R:Tsujiura, A.; Yasojima, K.; Hashimoto-Gotoh, T.

Biochem. Biophys. Res. Commun. 231, 392-396, 1997
A:Title: Cloning of Xenopus presenilin-alpha and -beta cDNAs and their differential e
A:Reference number: JC5390; MUID:97223465
A:Accession: JC5390
A>Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-433 <TSU>
A:Cross-references: DBJ:D84427; NID:gl944353; PIDN:BAAL9570.1; PID:dl020347; PID:gl9
A:Experimental source: brain
C:Comment: This protein plays a role in negative regulation of apoptotic cascades dur
C:Superfamily: presenilin
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F:99-119/Domain: transmembrane #status predicted <TM2>
F:130-149/Domain: transmembrane #status predicted <TM3>
F:161-178/Domain: transmembrane #status predicted <TM4>
F:187-203/Domain: transmembrane #status predicted <TM5>
F:210-227/Domain: transmembrane #status predicted <TM6>
F:374-394/Domain: transmembrane #status predicted <TM7>
Query Match 74.4%; Score 1431.5; DB 2; Length 433;
Best Local Similarity 72.0%; Pred. No. 4.7e-101;
Matches 286; Conservative 27; Mismatches 59; Indels 25; Gaps 4;
QY 1 BELTLKYGAKHVMFLFVPVTLICMIVVATIKSVRYTEKNGQLIYTPPTEDTPSVGQRL 60
Db 37 BELTLKYGAKHVMFLFVPVTLICMIVVATIKSVSYTRFDGQLIYTPPTEDTESVGQRL 96
QY 61 NSVLNTLMISVIVVMTIFLVLYKYCYKFIHGWLMISLMFLFYIYLGVEVLTYN 120
Db 97 NSILNATMISVIVMTILLVLYKYCYKFIHGWLMISLSSLLLFYFIFGLVEVFTYN 156
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Db 157 VADYVTVALLIWNFGVGMICIHKGKPLLOQAYLIMISALMALVFIKYLPEWTWLLIL 216
QY 181 GAISYDVLAVLCPKGLRMLVETAQERNEPIFPALIIYSAMVTVGMKLDPS-SQGL 239
Db 217 AVISYDVLAVLSPKGLRMLVETAQERNETLFPALIIYSSTMIWLVNMAADGDFGLKQAS 276
QY 240 QLPYDPEN-----EDSDSFCEPSYVEFP-----PLGYGPEEL----- 277
Db 277 TKYNTQAPTAPHRSDSAASDDNGGFDTHEDHNAQGPINSTPESRVAVQALPNSPP 336
QY 277 ---EEERGVKLGIDFIFYSVLVGKAAATGSGDWNNTLACFAVAILGICLTLTLLAVFK 335
Db 337 SEDPERGVKLGIDFIFYSVLVGKASATASGDWNTTACFAVAILGICLTLTLLAIFK 396
QY 336 ALPALPISITIFGLIFYSTDNLRPFMDTLASHQLYI 372
Db 397 ALPALPISITIFGLVFFATDYLVPFMDQLAFHQFYI 433
RESULT 7
S58396
presenilin 1, splice form 467 - human
N:Alternate names: Alzheimer's disease protein 3; protein S182
C:Species: Homo sapiens (man)
C:Date: 29-Jan-1998 #sequence_revision 13-Feb-1998 #text_change 29-Sep-1999
C:Accession: S58396; S71401; S71402
R:Sherrington, R.; Rogaev, E.I.; Liang, Y.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; C
ero, I.; Pinessi, L.; Nee, L.; Chumakov, I.; Pollen, D.; Brookes, A.; Sanseau, P.; Po
E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 375, 754-760, 1995
A:Title: Cloning of a gene bearing missense mutations in early-onset familial Alzheim
A:Reference number: I58095; MUID:95319502
A:Accession: S58396
A:Molecule type: mRNA
A:Residues: 1-467 <SHE>
A:Cross-references: EMBL:L42110; NID:g904118; PIDN:AAB46416.1; PID:g904119
R:Vidal, R.; Ghiso, J.; Wisniewski, T.; Frangione, B.
FEBS Lett. 393, 19-23, 1996
A:Title: Alzheimer's presenilin 1 gene expression in platelets and megakaryocytes. Id

[illegible]

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Qy 181 GAISYDLDVAVLCPKGPLRMLVETAQERNPEIFPALLIYSSAMVWTVGMALDPSQCAL- 240
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 251 AVISYDLDVAVLCPKGPLRMLVETAQERNPEIFPALLIYSSAMVWTVGMALDPSQCAL- 310
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Qy 240 -QLPYD-----PEMEEDSYDSFGPEPSYPEFEP-----PLTGYGCEEL----- 277
      || : : : : : || : : : : : || : : : : : || : : : : : || : : : : :
Db 311 KNTYKNAQGTREAAQAVPNDGCGFSEWEAQRDSQLGHRSTSVSRAAVOEISSIP 370
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Qy 277 -EEEEERGVKLGIDGFYFYSVLGKAAATSGDWNNTTLACFVAILIGLCITLLLLAVFKK 335
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 371 SEDPEERGVKLGIDGFYFYSVLGKASATASGDWNNTTLACFVAILIGLCITLLLLAIFKK 430
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Qy 336 ALPALPISITFGLIFYSTDNLVRFPMDTLASHOLYI 372
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 431 ALPALPISITFGLVFYFATDYLQVPMQLAFHQFYI 467
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 11
S63684
presenilin 1, splice form 374 - human
N:Alternate names: Alzheimer's disease protein 3
C:Species: Homo sapiens (man)
C>Date: 20-Jul-1996 #sequence_revision 13-Mar-1997 #text_change 29-Sep-1999
C:Accession: S63684
R:Sahara, N.; Yahagi, Y.; Takagi, H.; Kondo, T.; Okochi, M.; Usami, M.; Shirasawa, T.
FEBS Lett. 381, 7-11, 1996
A:Title: Identification and characterization of presenilin I-467, I-463 and I-374.
A:Reference number: S63683; MUID:96193901
A:Accession: S63684
A>Status: preliminary; nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-374 <SAH>
A:Cross-references: EMBL:U40380; NID:g1244639; PIDN:AAB05895.1; PID:g1244640
C:Genetics:
A:Gene: GDB:PSEN1; AD3; FAD; S182; P51
A:Cross-references: GDB:135682; OMIM:104311
A:Map position: 14q24.3-14q24.3
C:Superfamily: presenilin

Query Match 52.6%; Score 1011; DB 2; Length 374;
Best Local Similarity 81.8%; Pred. No. 2.4e-69;
Matches 193; Conservative 18; Mismatches 25; Indels 0; Gaps 0;

Qy 1 BELTLKYGAKHVMILFVPVTLICMIVVATIKSVFYETKNGQLIYTPFTEDTPSVGORLL 60
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 67 BELTLKYGAKHVMILFVPVTLICMIVVATIKSVFYETKNGQLIYTPFTEDTPSVGORAL 126
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Qy 61 NSVLNTLMISIVVMVTFVLVLYKYRCYFIHGWLMSSLMLLFLFTYIYLGVEVLKTYN 120
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 127 HSILNAALMISIVVMVILLVLYKYRCYKVIHAWLISSLLLLFFSFYILGVEVFTYN 186
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Qy 121 VANDYPTLLTVNFGAVGVCVHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAVVIL 180
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 187 VADVYITVALLIWNFGVVGMSIHWKGPLRLOQAYLIMISALMALVFIKYLPEWTAWIL 246
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Qy 181 GAISYDLDVAVLCPKGPLRMLVETAQERNPEIFPALLIYSSAMVWTVGMALDPSQ 236
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 247 AVISYDLDVAVLCPKGPLRMLVETAQERNPEIFPALLIYSSAMVWTVGMALDPSQAQ 302
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 12
S60253
sel-12 protein - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C>Date: 10-Apr-1996 #sequence_revision 19-Apr-1996 #text_change 13-Sep-1998
C:Accession: S60253
R:Levitan, D.; Greenwald, I.
Nature 377, 351-354, 1995
A:Title: Facilitation of lin-12-mediated signalling by sel-12, a Caenorhabditis elegans
A:Reference number: S60253; MUID:96032531
A:Accession: S60253
A>Status: preliminary; nucleic acid sequence not shown

```

|||||
Db 369 KGLGDEIFYSVLGRAAMY---DLMTVYACYLATISGLGCTLLILSVYNRALPALPISI 425
QY 345 TFGILFYFSTDNLVRPFM 362
Db 426 MLGVVYFYFLTRLLMEPFV 443
RESULT 15
A43459
sperm membrane protein spe-4 - Caenorhabditis elegans
N:Alternate names: probable integral membrane protein
C:Species: Caenorhabditis elegans
C>Date: 10-Jun-1993 #sequence:revision 18-Nov-1994 #text_change 09-Sep-1997
C:Accession: A43459; S24632; S24633
R:L'Hernault, S.W.; Arduengo, P.M.
J. Cell Biol. 119, 55-68, 1992
A:Title: Mutation of a putative sperm membrane protein in Caenorhabditis elegans prevents
A:Reference number: A43459; MUID:92407040
A:Accession: A43459
A>Status: preliminary; not compared with conceptual translation
A:Molecule type: DNA; mRNA
A:Residues: 1-465 <LHE>
A:Cross-references: EMBL:Z14067; NID:g6868; PID:g6869; EMBL:Z14066; NID:g6870; PID:g6871
A:Experimental source: strain Bristol N2
A:Note: the nucleotide sequence was submitted to the EMBL Data Library, July 1992
C:Genetics:
A:Introns: 69/3; 154/3; 200/1; 224/3; 300/1; 386/1; 435/1

Query Match 14.28; Score 274; DB 2; Length 465;
Best Local Similarity 21.58; Pred. No. 1.5e-13;
Matches 93; Conservative 80; Mismatches 127; Indels 132; Gaps 14;
QY 38 EKNQGLIYPTFEDT--PSVGORLINSVLN---TLIMISIVVMTIFLVLYKRCYKF 91
Db 42 EVNSELSKTYFLDPSFEQTTGNLLDGFINGVGTILVGCVSFIMLAF--VLFDER--RI 97
QY 92 IHGWLIMSSMLLF-----LFTYIYLGVLKTYNVANDYPTLLL-----TWNFGA 137
Db 98 VKAWLTLSCLLILFGVSAQTLHDMFSQVFDQDDNNQY-----YMTIVLIIVPTVVYGF- 152
QY 138 VGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWILGAISVYDLVAVLCPKGP 197
Db 152 -GIYAFPSNSSLILHQIFVVTNCSLIISVYLKRVFPKTFWFLWVFLFVLDLFAVLAPMGP 210
QY 198 LRMIVETAQERNEPIFPALIIYSSAMVTVGMKLDPSQG-----ALQLPYDPE 246
Db 211 LKKVOEKASDYKCVNLIMFSANEKRLTAGSNEETNEGEESTIRRVKOTIEYTTRE 270
QY 247 MEEDSY-----DSF-----GEPSTYFVFPPLTGYPGEELEEE--- 282
Db 271 AQDDFYQKIRQRAAINPDSVPTSEHSPLEAEPSPIELKEKNST-----EELSDDESOTS 326
QY 282 -----R 282
Db 327 ETSSGNSLSSDSTTVSTSDISTAECDQKWDLVNSLNPNNDKRPATAADALNDGE 386
QY 283 GVKLGLGDFIFYSLVGVKAAATGSDWNTTLACFVAIILGLCLTLALLAVFKKALPALPI 342
Db 387 VLRIGFGDFVYSLIGQAASGCP--FAVISALGILFLGLVVLTVTFSTEESTPALPL 444
QY 343 SITEGLIFYFST 354
Db 445 PVICGTFCYFSS 456

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 18:09:18 ; Search time 32.57 seconds
(without alignments)
341.103 Million cell updates/sec

Title: US-08-509-359B-138
Perfect score: 1923
Sequence: 1 EELTLAYGAKHVMFLPVPY.....STDNLVRFPMDTLASHQLYI 372

Scoring table: BLOSUM62

Searched: 82229 seqs, 29864866 residues

Database : SwissProt_38.*

Word size : 0

Number of hits that pass the threshold : 82229

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1923	100.0	448	1	PSN2_HUMAN
2	1876	97.6	448	1	PSN2_MOUSE
3	1866	97.0	448	1	PSN2_RAT
4	1853	96.5	445	1	PSN2_MICMU
5	1534	79.8	449	1	PSN2_XENLA
6	1431.5	74.4	467	1	PSN1_MOUSE
7	1431.5	74.4	433	1	PSN1_XENLA
8	1431	74.4	468	1	PSN1_RAT
9	1429.5	74.3	467	1	PSN1_HUMAN
10	1413.5	73.5	467	1	PSN1_MICMU
11	1127.5	58.6	541	1	PSN1_DROME
12	1032	53.7	836	1	YL4K_CAEEL
13	972	50.5	461	1	SE12_CAEEL
14	524.5	27.3	358	1	HOP1_CAEEL
15	513	26.7	433	1	PSNH_ARATH
16	274	14.2	465	1	SPE4_CAEEL
17	110	5.7	2016	1	CIN5_HUMAN
18	108	5.6	381	1	CYB_DASCR
19	103	5.4	382	1	CYB_DIDMA
20	103	5.4	380	1	CYB_MICLO
21	102.5	5.3	381	1	CYB_ANTFL
22	102	5.3	381	1	CYB_DASMA
23	102	5.3	381	1	CYB_PSENI
24	101.5	5.3	381	1	CYB_NINIV
25	101.5	5.3	381	1	CYB_PAPAP
26	101	5.3	381	1	CYB_PLAMS
27	100.5	5.2	381	1	CYB_PSEMD
28	100	5.2	381	1	CYB_DASGE
29	99.5	5.2	1840	1	CIN4_RAT
30	99	5.1	2019	1	CIN5_RAT
31	99	5.1	381	1	CYB_ANTMI
32	99	5.1	381	1	CYB_PHATA
33	98.5	5.1	331	1	YDFG_SCHPO
34	98	5.1	381	1	CYB_ANTME
35	98	5.1	381	1	CYB_DASAL
36	98	5.1	381	1	CYB_SMICR
37	98	5.1	460	1	NU4M_GADMO
38	98	5.1	217	1	YPR4_ECOLI
39	97.5	5.1	381	1	CYB_DASHA

40 97.5 5.1 381 1 CYB_PARBI Q35380 parantechin
41 97 5.0 238 1 ATP6_BACP3 P09218 bacillus ps
42 97 5.0 381 1 CYB_DASYI Q34399 dasyurus vi
43 97 5.0 381 1 CYB_MACRO P92671 macropus ro
44 97 5.0 347 1 ETBR_COTJA Q90328 coturnix co
45 97 5.0 447 1 NU4M_APILI P34853 apis mellif

ALIGNMENTS

RESULT 1
PSN2_HUMAN
ID PSN2_HUMAN STANDARD; PRT; 448 AA.
AC P49810;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN 2 (PS-2) (STM-2) (E5-1) (AD3LP) (AD5).
GN PSN2 OR PSN12 OR AD4 OR PS2 OR STM2.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A., AND VARIANT FAD ILE-141.
RX MEDLINE; 95355816.
RA LEVI-LAHAD E., WASCO W., POORRAJ P., ROMANO D.M., OSHIMA J.,
RA PETTINGELL W.H., YU C.-E., JONDRO P.D., SCHMIDT S.D., WANG K.,
RA CROWLEY A.C., FU Y.-H., GUENETTE S.Y., GALAS D., NEMENS E.,
RA WIJSMAN E.M., BIRD T.D., SCHELLENBERG G.D., TANZI R.E.;
RT "Candidate gene for the chromosome 1 familial Alzheimer's disease
RT locus.";
RL Science 269:973-977(1995).
RN [2]
RP SEQUENCE FROM N.A., AND VARIANTS FAD ILE-141 AND VAL-239.
RC TISSUE-BRAIN, AND COLON;
RX MEDLINE; 95379971.
RA ROGAEV E.I., SHERRINGTON R., ROGAEVA E.A., LEVESQUE G., IKEDA M.,
RA LIANG Y., CHI H., LIN C., HOLMAN K., TSUDA T., MAR L., SORBI S.,
RA NACMIAS B., PIACENTINI S., AMADUGGI L., CHUMAKOV I., COHEN D.,
RA LANNFELT L., FRASER P.E., ROMMENS J.M., ST GEORGE-HYSLOP P.H.;
RT "Familial Alzheimer's disease in kindreds with missense mutations in
RT a gene on chromosome 1 related to the Alzheimer's disease type 3
RT gene.";
RL Nature 376:775-778(1995).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE; 96109229.
RA LI J., MA J., POTTER H.;
RT "Identification and expression analysis of a potential familial
RT Alzheimer disease gene on chromosome 1 related to AD3.";
RL Proc. Natl. Acad. Sci. U.S.A. 92:12180-12184(1995).
RN [4]
RP SEQUENCE FROM N.A.
RA LEVI-LAHAD E., POORRAJ P., WANG K., FU Y.-H., OSHIMA J.,
RA MULLIGAN J., SCHELLENBERG G.D.;
RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
RN [5]
RP REVIEW ON VARIANTS.
RX MEDLINE; 98180715.
RA CRUTS M., VAN BROECKHOVEN C.;
RT "Presenilin mutations in Alzheimer's disease.";
RL Hum. Mutat. 11:183-190(1998).
RN [6]
RP VARIANT AD HIS-62.
RX MEDLINE; 98046005.
RA CRUTS M., VAN DUIN C.M., BACKHOVENS H., VAN DEN BROECK M.,
RA WEHNET A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J.,
RA ST GEORGE-HYSLOP P.H., HOFMAN A., VAN BROECKHOVEN C.;
RT "Estimation of the genetic contribution of presenilin-1 and -2
RT mutations in a population-based study of presenile Alzheimer
RT disease.";
RL Hum. Mol. Genet. 7:43-51(1998).

DR PFAM; PF01080; Presenilin; 1.
KW Transmembrane; Alternative Initiation.
FT CHAIN 1 448 PRESENILIN 2.
FT CHAIN 298 448 PRESENILIN 2-SHORT.
FT TRANSNEM 88 106 POTENTIAL.
FT TRANSNEM 142 160 POTENTIAL.
FT TRANSNEM 167 188 POTENTIAL.
FT TRANSNEM 203 219 POTENTIAL.
FT TRANSNEM 230 246 POTENTIAL.
FT TRANSNEM 253 269 POTENTIAL.
FT TRANSNEM 288 305 POTENTIAL.
FT TRANSNEM 317 406 POTENTIAL.
FT TRANSNEM 413 429 POTENTIAL.
FT CONFLICT 87 87 R -> H (IN REF. 2).
FT CONFLICT 226 226 R -> V (IN REF. 2).
FT CONFLICT 324 324 MISSING (IN REF. 2).
SQ SEQUENCE 448 AA; 49955 MW; 680ACF19 CRC32;

Query Match 97.6%; Score 1876; DB 1; Length 448;
Best Local Similarity 97.3%; Pred. No. 1.7e-121;
Matches 362; Conservative 3; Mismatches 7; Indels 0; Gaps 0;

QY 1 BELTLKYGAKHVMFLFVPTLCMIVVATIKSVRFYTEKNGQLIYPTFTEDTPSGVQRLL 60
DB 77 BELTLKYGAKHVMFLFVPTLCMIVVATIKSVRFYTEKNGQLIYPTFTEDTPSGVQRLL 136

QY 61 NSVLNTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSLMFLFTYIYLGVLKTYN 120
DB 137 NSVLNTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSLMFLFTYIYLGVLKTYN 196

QY 121 VAMDYPTLLTWNFGAVGWCIIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWIL 180
DB 197 VAMDYPTLLTWNFGAVGWCIIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWIL 256

QY 181 GAISYDVLAVLCPKGPLRMLVETAQERNEPFPALIISSAMVWTVGMKLDPSQGAQ 240
DB 257 GAISYDVLAVLCPKGPLRMLVETAQERNEPFPALIISSAMVWTVGMKLDPSQGAQ 316

QY 241 LPYDPEMEEDSYDSFGSPSYPEVPEPLTGYPGEELEEEERGVKLGDFIFYSVLVGK 300
DB 317 LPYDPEMEEDSYDSFGSPSYPEVPEPLTGYPGEELEEEERGVKLGDFIFYSVLVGK 376

QY 301 AAATGSGDWNNTLACFAVAILIGLCTLLLLAVFKKALPALPISITFGLIFFYFSTDNLVRP 360
DB 377 AAATGSGDWNNTLACFAVAILIGLCTLLLLAVFKKALPALPISITFGLIFFYFSTDNLVRP 436

QY 361 FMDTLASHQLYI 372
DB 437 FMDTLASHQLYI 448

RESULT 3
PSN2_RAT STANDARD; PRT; 448 AA.
AC O88777; O35546; O08947;
DT 13-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN 2 (PS-2).
GN PSN2 OR PSN2 OR PS2.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-WISTAR; TISSUE-BRAIN;
RA FRENZEL S., ABDEL A.S., LUEBBERT H.;
RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-WISTAR; TISSUE-BRAIN;
RX MEDLINE; 97473536.

RA TAKAHASHI H., MERCKEN M., NAKAZATO Y., NOGUCHI K., MURAYAMA M.,
RA IMAHORI K., TAKASHIMA A.;
RT "Cloning of cDNA and expression of the gene encoding rat
RT presenilin-2.";
RL Gene 197:383-387(1997).
RN [3]
RN SEQUENCE FROM N.A.
RC STRAIN-WISTAR; TISSUE-BRAIN;
RX MEDLINE; 98207716.
RA TAKAHASHI H., TABIRA T.;
RT "Cloning of the cDNA encoding rat presenilin-2.";
RL Biochim. Biophys. Acta 1396:259-262(1998).
CC -1- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE
CC EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE. MAY
CC FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS (BY
CC SIMILARITY).
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
CC -----
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CC -----
CC EMBL; X99267; CAA67663.1; -;
DR EMBL; D83700; BAA22832.1; -;
DR EMBL; AB004454; BAA20406.1; -;
DR PFAM; PF01080; Presenilin; 1.
KW Transmembrane.
FT TRANSNEM 88 106 POTENTIAL.
FT TRANSNEM 142 160 POTENTIAL.
FT TRANSNEM 167 188 POTENTIAL.
FT TRANSNEM 203 219 POTENTIAL.
FT TRANSNEM 230 246 POTENTIAL.
FT TRANSNEM 253 269 POTENTIAL.
FT TRANSNEM 288 305 POTENTIAL.
FT TRANSNEM 387 406 POTENTIAL.
FT TRANSNEM 413 429 POTENTIAL.
FT CONFLICT 7 7 S -> T (IN REF. 1).
FT CONFLICT 86 87 KH -> ND (IN REF. 3).
SQ SEQUENCE 448 AA; 50051 MW; E0DF681 CRC32;

Query Match 97.0%; Score 1866; DB 1; Length 448;
Best Local Similarity 96.8%; Pred. No. 8.4e-121;
Matches 360; Conservative 4; Mismatches 8; Indels 0; Gaps 0;

QY 1 BELTLKYGAKHVMFLFVPTLCMIVVATIKSVRFYTEKNGQLIYPTFTEDTPSGVQRLL 60
DB 77 BELTLKYGAKHVMFLFVPTLCMIVVATIKSVRFYTEKNGQLIYPTFTEDTPSGVQRLL 136

QY 61 NSVLNTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSLMFLFTYIYLGVLKTYN 120
DB 137 NSVLNTLMISVIVMTIFLVLYKYRCYKFIHGWLMSSLMFLFTYIYLGVLKTYN 196

QY 121 VAMDYPTLLTWNFGAVGWCIIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWIL 180
DB 197 VAMDYPTLLTWNFGAVGWCIIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWIL 256

QY 181 GAISYDVLAVLCPKGPLRMLVETAQERNEPFPALIISSAMVWTVGMKLDPSQGAQ 240
DB 257 GAISYDVLAVLCPKGPLRMLVETAQERNEPFPALIISSAMVWTVGMKLDPSQGAQ 316

QY 241 LPYDPEMEEDSYDSFGSPSYPEVPEPLTGYPGEELEEEERGVKLGDFIFYSVLVGK 300
DB 317 LPYDPEMEEDSYDSFGSPSYPEVPEPLTGYPGEELEEEERGVKLGDFIFYSVLVGK 376

QY 301 AAATGSGDWNNTLACFAVAILIGLCTLLLLAVFKKALPALPISITFGLIFFYFSTDNLVRP 360
DB 377 AAATGSGDWNNTLACFAVAILIGLCTLLLLAVFKKALPALPISITFGLIFFYFSTDNLVRP 436

Db 191 VAVDYITVALLIWFVGVGMIAHWRKPLRLQAOYLIMISALMALVFIKYLPEWTAWLIL 250
QY 181 GATSVYDLVAVLCCKPLRLMVLTAQERNEPFPALITYSSAMVTWGMKLDPSOGALQ 240
Db 251 AVISYDLVAVLCCKPLRLMVLTAQERNEPFPALITYSSAMVTWGMKLDPSOGALQ--RR 308
QY 241 LPYDP-----EMEEDSYSGFSPSPYEPFPLTGYG-----EEL--- 277
Db 309 VPKNPYSTGOTERETQDTGTGSDGGSEWEAQRDHLGPHRSTPSRAVQELSGS 368
QY 277 -----EDEBERGVKLGDFIFYSVLVKGAAATGSGDWNTTACFAVAILGLCUTLLAV 332
Db 369 ILTSEDEPBERGVKLGDFIFYSVLVKGAAATGSGDWNTTACFAVAILGLCUTLLAV 428
QY 333 FKALPALPISITGFLIFYSFDTNLRVPRMDTLASHQLYI 372
Db 429 FKALPALPISITGFLIFYSFDTNLRVPRMDTLASHQLYI 468
RESULT 9
PSN1_HUMAN
ID PSN1_HUMAN STANDARD; PRT: 467 AA.
AC P49768; Q14762; Q15719; Q15720;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN 1 (PS-1) (S182 PROTEIN).
GN PSN1 OR PSN1 OR A3 OR PSI.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.; AND VARIANTS AD (FORMS I-467 AND I-463).
RC TISSUE=BRAIN;
RX MEDLINE; 95319502.
RA SHERRINGTON R., ROGAJEV E.I., LIANG Y., ROGAJEV E.A., LEVESQUE G.,
RA IKEDA M., CHI H., LIN C., HOLMAN K., TSUDA T., MAR L.,
RA FONGIN J.-F., BRUNI A.C., MONTESI M.P., SORBI S., RAINERO I.,
RA PINESSI L., NEE L., CHUMAKOV I., POLLEN D., BROOKS A.,
RA SANSEAU P., POLINSKY R.J., WASSO W., DA SILVA H.A.R., HAINES J.L.,
RA PERICAK-VANCE M.A., TANZI R.E., ROSES A.D., FRASER P.E.,
RA ROMMENS J.M., ST GEORGE-HYSLOP P.H.;
RT "Cloning of a gene bearing missense mutations in early-onset familial
RT Alzheimer's disease.";
RL Nature 375:754-760(1995).
RN [2]
RP SEQUENCE FROM N.A. (FORMS I-463 AND I-374).
RC TISSUE=BLOOD, AND BRAIN;
RX MEDLINE; 96193901.
RA SAHARA N., YAHAGI Y.-I., TAKAGI H., KONDO T., OKOCHI M., USAMI M.,
RA SHIRASAWA T., MORI H.;
RT "Identification and characterization of presenilin I-463 and
RT I-374.";
RL FEBS Lett. 381:7-11(1996).
RN [3]
RP SEQUENCE OF 1-113 FROM N.A.
RA TSUJIMURA A., HASHIMOTO-GOTOH T.;
RL Submitted (MAR-1996) to the EMBL/GenBank/DBJ databases.
RN [4]
RP REVIEW ON VARIANTS.
RX MEDLINE; 98180715.
RA CRUTS M., VAN BROECKHOVEN C.;
RT "Presenilin mutations in Alzheimer's disease.";
RL Hum. Mutat. 11:183-190(1998).
RN [5]
RP VARIANTS AD THR-143 AND ALA-384.
RX MEDLINE; 96177673.
RA CRUTS M., BACKHOVENS H., WANG S.-Y., VAN GASSEN G., THEUNS J.,
RA DE JONGHE C., WEHRT A., DE VOECHT J., DE WINTER G., CRAS P.,
RA BRUYLAND M., DATSON N., WEISSENBAACH J., DEN DUNNEN J.T., MARTIN J.-J.,
RA HENDRIKS L., VAN BROECKHOVEN C.;
RT "Molecular genetic analysis of familial early-onset Alzheimer's
RT disease linked to chromosome 14q24.3.";

Hum. Mol. Genet. 4:2363-2372(1995).
RN [6]
RP VARIANTS AD L-82; H-115; T-139; R-163; T-231; L-264; V-392 AND Y-410.
RX MEDLINE; 96177674.
RA CAMPION D., FLAMAN J.-M., BRICE A., HANNEQUIN D., DUBOIS B.,
RA MARTIN C., MOREAU V., CHARBONNIER F., DIDIERJEAN O., TARDIEU S.,
RA PENET C., PUEL M., PASQUET F., LE DOZE F., BELLIS G., CALENDA A.,
RA HEILIG R., MARTINEZ M., MALLET J., BELLIS M., CLERGET-DARPOUX F.,
RA AGID Y., FREBOURG T.;
RT "Mutations of the presenilin I gene in families with early-onset
RT Alzheimer's disease.";
RL Hum. Mol. Genet. 4:2373-2377(1995).
RN [7]
RP VARIANTS AD VAL-260; VAL-285 AND VAL-392.
RX MEDLINE; 95379971.
RA ROGAJEV E.I., SHERRINGTON R., ROGAJEV E.A., LEVESQUE G., IKEDA M.,
RA LIANG Y., CHI H., LIN C., HOLMAN K., TSUDA T., MAR L., SORBI S.,
RA NACMIAS B., PIACENTINI S., AMADUCCI L., CHUMAKOV I., COHEN D.,
RA LANFELT L., FRASER P.E., ROMMENS J.M., ST GEORGE-HYSLOP P.H.;
RT "Familial Alzheimer's disease in kindreds with missense mutations in
RT a gene on chromosome 1 related to the Alzheimer's disease type 3
RT gene.";
RL Nature 376:775-778(1995).
RN [8]
RP VARIANTS AD V-139; V-146; Y-163; T-267; A-280 AND G-280.
RX MEDLINE; 96024664.
RA CLARK R.F., HUTTON M., FULDNER R.A., FROELICH S., KARRAN E.,
RA TALBOT C., CROOK R., LENDON C., PRIHAR G., HE C., KORENBLAT K.,
RA MARTINEZ A., WRAGG M., BUSFIELD F., BEHRENS M.I., MYERS A., NORTON J.,
RA MORRIS J., MEHTA N., PEARSON C., LINCOLN S., BAKER M., DUFF K.,
RA ZEHRE C., PEREZ-TUR J., HOULDEN H., RUIZ A., OSSA J., LOPERA F.,
RA ARCOS M., MADRIGAL L., COLLINGE J., HUMPHREYS C., ASWORTH T.,
RA SARNER S., FOX N., HARVEY R., KENNEDY A., ROQUES P., CLINE R.T.,
RA PHILLIPS C.A., VENTER J.C., FORSEL L., AXELMAN K., LILIUS L.,
RA JOHNSTON J., COMBURN R., VIITANEN M., WINBLAD B., KOSIK K., HALTIA M.,
RA POYHONEN M., DICKSON D., MANN D., NEARY D., SNOWDEN J., LANTOS P.,
RA LANFELT L., ROSSOR M., ROBERTS G.W., ADAMS M.D., HARDY J., GOATE A.,
RT "The structure of the presenilin 1 (S182) gene and identification of
RT six novel mutations in early onset AD families. Alzheimer's Disease
RT Collaborative Group.";
RL Nat. Genet. 11:219-222(1995).
RN [9]
RP VARIANTS AD PHE-96; ARG-163 AND THR-213.
RX MEDLINE; 96310408.
RA KAMINO K., SATO S., SAKAKI Y., YOSHIWA A., NISHIWAKI Y., TAKEDA H.,
RA TANABE H., NISHIMURA T., LI K., ST GEORGE-HYSLOP P.H., MIKI T.,
RA OGIHARA T.;
RT "Three different mutations of presenilin 1 gene in early-onset
RT Alzheimer's disease families.";
RL Neurosci. Lett. 208:195-198(1996).
RN [10]
RP VARIANT AD ALA-280.
RX MEDLINE; 97442268.
RA LENDON C.L., MARTINEZ A., BEHRENS I.M., KOSIK K.S., MADRIGAL L.,
RA NORTON J., NEUMAN R., MYERS A., BUSFIELD F., WRAGG M., ARCOS M.,
RA ARANGO VIANA J.C., OSSA J., RUIZ A., GOATE A.M., LOPERA F.;
RT "E280A PS-1 mutation causes Alzheimer's disease but age of onset is
RT not modified by ApoE alleles.";
RL Hum. Mutat. 10:186-195(1997).
RN [11]
RP VARIANTS AD THR-233 AND THR-278.
RX MEDLINE; 97316242.
RA KWOK J.B.J., TADDEI K., HALLUPP M., FISHER C., BROOKS W.S., BROE G.A.,
RA HARDY J., FULHAM M.J., NICHOLSON G.A., STELL R.,
RA ST GEORGE-HYSLOP P.H., FRASER P.E., KAKULAS B., CLARINETTE R.,
RA RELIN N., GANDY S.E., SCHOFIELD P.R., MARTINS R.N.;
RT "Two novel (M233T and R278T) presenilin-1 mutations in early-onset
RT Alzheimer's disease pedigrees and preliminary evidence for
RL association of presenilin-1 mutations with a novel phenotype.";
RN NeuroReport 8:1537-1542(1997).
RN [12]
RP VARIANT GLY-318.
RX MEDLINE; 99115106.

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CC -----

DR EMBL; Z71333; CAA95930.1; -;
DR PFAM; PF01080; Presenilin; 1.
KW Transmembrane; Alternative splicing; Glycoprotein.
FT TRANSMEM 83 103 POTENTIAL.
FT TRANSMEM 133 153 POTENTIAL.
FT TRANSMEM 161 181 POTENTIAL.
FT TRANSMEM 191 211 POTENTIAL.
FT TRANSMEM 221 241 POTENTIAL.
FT TRANSMEM 244 264 POTENTIAL.
FT TRANSMEM 281 301 POTENTIAL.
FT TRANSMEM 408 428 POTENTIAL.
FT TRANSMEM 433 453 POTENTIAL.
FT CARBOHYD 279 279 POTENTIAL.
FT CARBOHYD 405 405 POTENTIAL.
FT VARSPPLIC 26 29 MISSING (IN ISOFORM I-463).
SQ SEQUENCE 467 AA; 52384 MW; A841A0B7 CRC32;

Query Match 73.5%; Score 1413.5; DB 1; Length 467;
Best Local Similarity 71.0%; Pred. No. 7.2e-90;
Matches 282; Conservative 34; Mismatches 56; Indels 25; Gaps 4;

Qy 1 BELTLKYGAKHVMLEFVPTLCMIVVATIKSVRYTEKNGOLIYPTPTDTPSVGQRL 60
Db 71 BELTLKYGAKHVMLEFVPTLCMIVVATIKSVRYTEKNGOLIYPTPTDTPSVGQRL 130
Qy 61 NSVLNTLMISVIVVMTIFLVLYKRYCYKFTGHWLIMSSLMMLFLFYIYLGVLKYN 120
Db 131 HSLVNAALMISVIVVMTIFLVLYKRYCYKFTGHWLIMSSLMMLFLFYIYLGVLKYN 190
Qy 121 VAMDYPTLLTWNFGAVGMCVCIHKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVIL 180
Db 191 VAVDYITVALLIMNFGVVGMSIHWKGPLRLOQAYLIMISALMALVFIKYLPEWTAWLIL 250
Qy 181 GAISYDVLAVLCPKGPLRMLVETAEQNERIFFPALLYSSAMVTVGMAKLDPPSQGAL- 240
Db 251 AVISYDVLAVLCPKGPLRMLVETAEQNETLFPALLYSSAMVTVGMAKLDPPSQGAL- 310
Qy 240 -QLPYD-----PEMEEDSYDSFGSPSYPEVPEP-----PLTGYGPEEL----- 277
Db 311 KNTYNAQGTERRAQASVPENDDGGFSEWEAQRDSQGLPHRSVTSVRAAQQEISSIPA 370
Qy 277 -EEEEERGVKGLGDFIFYSVLVGKAATGSDWNTTILACFVAILIGLCLTLLLAFFKK 335
Db 371 SEDPEERGKVLGDFEYFVSVLVGKASATASGDWNTTILACFVAILIGLCLTLLLAFFKK 430
Qy 336 ALPALPISITGLFIYFTSDNLNRPDMTTLASHQYI 372
Db 431 ALPALPISITGLFIYFATDYLVOPFMDQLAFHQFYI 467

RESULT 11

ID PSN_DROME STANDARD; PRT; 541 AA.
AC 002194; 002395; 076802;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN HOMOLOG (DPS) (DMPS).
GN PS.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
RN [1]
RP SEQUENCE FROM N.A. (LONG ISOFORM).
RX MEDLINE; 97285868.
RA BOULIANNE G.L., LYNE-BAR I., HUMPHREYS J.M., LIANG Y., LIN C.,
RA ROGAEE E., ST GEORGE-HYSLOP P.;
RT "Cloning and characterization of the Drosophila presenilin

RT homologue.";
RL NeuroReport 8:1025-1029(1997).
RN [2]
RP SEQUENCE FROM N.A. (SHORT ISOFORM).
RC STRAIN=CANTON-S;
RA HONG C.S., KOO E.H.;
RT "Isolation and characterization of Drosophila presenilin homolog.";
RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A. (LONG AND SHORT ISOFORMS).
RA YE Y., FORTINI M.E.;
RT "Characterization of Drosophila presenilin and its colocalization
with Notch during development.";
RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
CC -----
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CC -----
DR EMBL; U77934; AAB61139.1; -;
DR EMBL; U78084; AAB53369.1; -;
DR EMBL; AF084184; AAC33129.1; -;
DR EMBL; AF084184; AAC33128.1; -;
DR FLYBASE; FBgn0019947; PS.
DR PFAM; PF01080; Presenilin; 1.
KW Transmembrane; Glycoprotein; Alternative splicing.
FT TRANSMEM 107 127 POTENTIAL.
FT TRANSMEM 155 175 POTENTIAL.
FT TRANSMEM 183 203 POTENTIAL.
FT TRANSMEM 217 237 POTENTIAL.
FT TRANSMEM 243 263 POTENTIAL.
FT TRANSMEM 266 286 POTENTIAL.
FT TRANSMEM 304 324 POTENTIAL.
FT TRANSMEM 482 502 POTENTIAL.
FT TRANSMEM 507 527 POTENTIAL.
FT CARBOHYD 129 129 POTENTIAL.
FT CARBOHYD 339 339 POTENTIAL.
FT CARBOHYD 410 410 POTENTIAL.
FT VARSPPLIC 384 397 MISSING (IN SHORT ISOFORM).
FT CONFLICT 80 81 GG -> RS (IN REF. 2).
SQ SEQUENCE 541 AA; 59304 MW; 796C4FE0 CRC32;

Query Match 58.6%; Score 1127.5; DB 1; Length 541;
Best Local Similarity 50.8%; Pred. No. 2.9e-70;
Matches 231; Conservative 61; Mismatches 74; Indels 89; Gaps 7;

Qy 1 BELTLKYGAKHVMLEFVPTLCMIVVATIKSVRYTEKNGOLIYPTPTDTPSVGQRL 60
Db 93 EEQGLKYGAQHVKLFVPSVLCMLVYVATINSIFYNSTDVLYLLYTPFHEQSPESVKFW 152
Qy 61 NSVLNTLMISVIVVMTIFLVLYKRYCYKFTGHWLIMSSLMMLFLFYIYLGVLKYN 120
Db 153 SALANSLIMSVVVMVTFLLIVLYKRCYRIHGMWILSSFMFLFTFYLYLEELRAYN 212
Qy 121 VAMDYPTLLTWNFGAVGMCVCIHKGPLVLOQAYLIMISALMALVFIKYLPEWSAWVIL 180
Db 213 IPMDYPTALLIMNFGVVGMSIHWKGPLRLOQAYLIFVAALMALVFIKYLPEWTAWVL 272
Qy 181 GAISYDVLAVLCPKGPLRMLVETAEQNERIFFPALLYSSAMVTV-----GMK 230
Db 273 AAISIWDLIAVLSPRGLRILVETAEQNEQIFPALLYSTVYVALVNTVTPQSOATAS 332
Qy 231 LDPSS-----QGALQLP-----YDPEMEEDSYDSFGSPSYPEV-FEP 266
Db 333 SSPSSNSTTTTTRATONSLASPEAAASQRTGNHPRONRDGGSVLATEGMPLVTFKS 392

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OM protein - protein search, using sw model

Run on: March 18, 2000, 22:07:43 ; Search time 49.26 Seconds
(without alignments)
523.595 Million cell updates/sec

Title: US-08-509-359B-138
Perfect score: 1923
Sequence: 1 EELTLKYGAKHVIMLFVPT.....STDLNRPFDMLASHQLYI 372

Scoring table: BLOSUM62

Searched: 225878 seqs, 69334122 residues

Database : SPTREMBL_12.*

Word size : 0

Number of hits that pass the threshold : 225878

- 1: sp.archaea.*
- 2: sp.bacteria.*
- 3: sp.fungi.*
- 4: sp.human.*
- 5: sp.invertebrate.*
- 6: sp.mammal.*
- 7: sp.mhc.*
- 8: sp.organelle.*
- 9: sp.phage.*
- 10: sp.plant.*
- 11: sp.rodent.*
- 12: sp.virus.*
- 13: sp.vertebrate.*
- 14: sp.unclassified.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
1	1896	98.6	449	6	Q9XT96	Q9xt96 bos taurus
2	1411.5	73.4	478	6	Q9XT97	Q9xt97 bos taurus
3	1398	72.7	456	13	Q9W6T7	Q9w6t7 brachydanio
4	1355	70.5	384	13	O73869	O73869 cyprinus ca
5	576.5	30.0	272	5	O96340	O96340 drosophila
6	402.5	20.9	184	4	O95465	O95465 homo sapien
7	113.5	5.9	406	5	Q19737	Q19737 caenorhabdi
8	112	5.8	458	13	Q42181	Q42181 fugu rubrip
9	110.5	5.7	320	8	Q34086	Q34086 coccyzus er
10	110.5	5.7	381	8	Q35425	Q35425 phascolosor
11	108.5	5.6	380	8	Q922C9	Q922c9 upupa epops
12	107.5	5.6	748	2	Q92577	Q92577 streptomyc
13	107	5.6	381	8	Q33723	Q33723 antechinus
14	106.5	5.5	382	8	Q34340	Q34340 didelphis m
15	106.5	5.5	318	11	P35294	P35294 rattus norv
16	106.5	5.5	303	11	P97829	P97829 rattus norv
17	106	5.5	652	5	Q93346	Q93346 caenorhabdi
18	106	5.5	382	8	Q34279	Q34279 didelphis a
19	106	5.5	382	8	Q34677	Q34677 glirophia ve
20	105	5.5	444	2	Q9X2N3	Q9x2n3 arthrobacte
21	105	5.5	379	8	Q34428	Q34428 echinys did
22	104	5.4	382	8	Q35561	Q35561 philander o
23	104	5.4	379	8	Q36096	Q36096 trinomys pa
24	104	5.4	379	8	Q34430	Q34430 echinys did
25	104	5.4	826	10	O80739	O80739 arabidopsis

ALIGNMENTS

RESULT 1

Q9XT96 PRELIMINARY; PRT; 449 AA.
AC Q9XT96;
DT 01-NOV-1999 (TREMBlrel. 12, Created)
DT 01-NOV-1999 (TREMBlrel. 12, Last sequence update)
DT 01-NOV-1999 (TREMBlrel. 12, Last annotation update)
DE PRESENILIN 2.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovinae; Bos.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=BRAIN;
RA SAHARA N., SHIRASAWA T., MORI H.;
RT "Molecular cloning of bovine presentin 2 gene."
RL Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF038937; AAD39024.1; -;
SQ SEQUENCE 449 AA; 50301 MW; A3DA878F CRC32;

Query Match	98.6%	Score	1896;	DB	6;	Length	449;
Best Local Similarity	98.4%	Pred.	No. 9.9e-131;				
Matches	366;	Conservative	2;	Mismatches	4;	Indels	0;
Gaps	0;						
Qy	1	EELTLKYGAKHVIMLFVPTLCMIVVATIKSVREYTEKNGOLIYTPFTEDTPSVGORLL	60				
Db	78	EELTLKYGAKHVIMLFVPTLCMIVVATIKSVREYTEKNGOLIYTPFTEDTPSVGORLL	137				
Qy	61	NSVLNTLMISVIVMTIFLVVLYKYRCYKFTGHWLIMSSMLLFLFYIYGEVLKTYN	120				
Db	138	NSVLNTLMISVIVMTIFLVVLYKYRCYKFTGHWLIMSSMLLFLFYIYGEVLKTYN	197				
Qy	121	VAMDYPTLLVWFGVGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLEWSAWVL	180				
Db	198	VAMDYPTLLVWFGVGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLEWSAWVL	257				
Qy	181	GAISYDVLAVLCPKGLRMLVETAQERNEFPFALIVSSAMVWTVGMKLPDSSOGALQ	240				
Db	258	GAISYDVLAVLCPKGLRMLVETAQERNEFPFALIVSSAMVWTVGMKLPDSSOGALQ	317				
Qy	241	LPYDPEMEEDSYDSFGSPYEPFPLTGYPGEELEEEERGVKLGDFIFYSVLVGK	300				
Db	318	LPYDPEMEEDSYDSFGSPYEPFPLTGYPGEELEEEERGVKLGDFIFYSVLVGK	377				
Qy	301	AAATGSGDNWTLACFAVILIGLCLTLLALLVFKKALPALPISITFGLIFYSTDNLRP	360				

Db 1 IMLFIPVTLKVVVVVATIKSVSYTKDQOQLIYPPFREDTETVQGRALNSMLNAAIMIS 60
Qy 72 VIVVMTIFLVLYKYRCYKFIHGLWLMSSMLLFTYIYLGELVKTNVAMDYPTLLLT 131
Db 61 VIVVMTLVVLYKYRCYKFIHGLWLMSSMLLFTYIYLGELVKTNVAMDYPTLLLT 120
Qy 132 VVNFAGVGVCHWKGPVLVQOAYLIMISALMALVFIKYLPEWSAWIIGALSIVYDLVAV 191
Db 121 IWNFGVGVCHWKGPVLVQOAYLIMISALMALVFIKYLPEWTAWLILAAISVYDLVAV 180
Qy 192 LCPKGPLRMLVETAQERNEPIFALIYSSAMVWVGMA-KLDPSOGALQLP----- 243
Db 181 LCPKGPLRILVETAQERNEPIFALIYSSAMVWVGMA-KLDPSOGALQLP----- 240
Qy 243 -----YDPMEDSDVSDFG-----EPSYDEVEPPLTGYPGEELEEEERGVKL 286
Db 241 APTAQPEDGGFTPAWVNOQHQGLQPMOSTEDSRIEIQELPSARPP--PVEDEERGVKL 298
Qy 287 GLGDFIFYSVLVGAATGSGDWNITLACFVAILIGLCLTLLLLAVFKKALPALPISITF 346
Db 299 GLGDFIFYSVLVGAATGSGDWNITLACFVAILIGLCLTLLLLAVFKKALPALPISITF 358
Qy 347 GLIFYFSTDLNVRPMDTLASHOLYI 372
Db 359 GLIFYFATDLNVRPMDQLAVHQFYI 384

RESULT 5
Qy 096340 PRELIMINARY; PRT; 272 AA.
AC 096340;
DT 01-MAY-1999 (TReMBLrel. 10, Created)
DT 01-MAY-1999 (TReMBLrel. 10, Last sequence update)
DT 01-MAY-1999 (TReMBLrel. 10, Last annotation update)
DE PRESENILIN (FRAGMENT).
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CANTON S;
RX MEDLINE; 98331525.
RA MARFANY G., DEL-FAVERO J., VALERO R., DE JONGHE C., WOODROW S.,
RA HENDRICKS L., VAN BROECKHOVEN C., GONZALEZ-DUARTE R.;
RT "Identification of a Drosophila presenilin homologue: evidence of
RT alternatively spliced forms";
RL J. Neurogenet. 12:41-54(1998).
DR EMBL; AF017025; AAD01611.1; .
FT NON_TER
SQ SEQUENCE 272 AA; 29456 MW; 606B9A5C CRC32;

Query Match 30.0%; Score 576.5; DB 5; Length 272;
Best Local Similarity 47.8%; Pred. No. 5.8e-35;
Matches 133; Conservative 29; Mismatches 41; Indels 75; Gaps 8;
Qy 164 ALVFIKYLPERSAWILGAISVYDLVAVLCPKGPLRMLVETAQERNEPIFALIYSSAMV 223
Db 1 ALVFIKYLPENTAWAVLAAISWDLVAVLSPRGLRILVETAQERNEQIFALIYSSIVV 60
Qy 224 WTV-----GMAKLDPS-----QGALQLPYDPEMEEDSDVSFGEPSYPE- 263
Db 61 YALVNTVTPQSOATASSPSSSSNTTTTRATQNSLA---SPEAAAASQRTGN-SHPRQ 116
Qy 263 -----VFEPLTGYPGE---ELEE----- 279
Db 117 NORDGSLVATEAEAGETQWSANLSERVARRQTEVOSTOSGNQORSNEYRTVTPDON 176
Qy 279 ----EEERGKVLGLGDFIFYSVLVGAATGSGDWNITLACFVAILIGLCLTLLLLAVFK 334
Db 177 HPDGOEERGKVLGLGDFIFYSVLVGAATGSGDWNITLACFVAILIGLCLTLLLLAVFK 234

Qy 335 KALPALPISITGLIFYFSTDLNVRPMDTLASHOLYI 372
Db 235 KALPALPISITGLIFYFSTDLNVRPMDTLASHOLYI 272
RESULT 6
Qy 095465 PRELIMINARY; PRT; 184 AA.
AC 095465;
DT 01-MAY-1999 (TReMBLrel. 10, Created)
DT 01-MAY-1999 (TReMBLrel. 10, Last sequence update)
DT 01-MAY-1999 (TReMBLrel. 10, Last annotation update)
DE MINILIN.
GN PSNI.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RA POWELL C.S., GEGG M.E., PALMER M.S.;
RT "Human presenilin 1 gene encodes an alternative protein-minilin.";
RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ008005; CAA07825.1; .
SQ SEQUENCE 184 AA; 21073 MW; 5C6FBAEE CRC32;

Query Match 20.9%; Score 402.5; DB 4; Length 184;
Best Local Similarity 73.5%; Pred. No. 2.1e-22;
Matches 83; Conservative 9; Mismatches 16; Indels 5; Gaps 1;
Qy 1 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFTEKNGQLIYPTFTDTPSVQRL 60
Db 71 EELTKYGAHVIMLFVPTLCMIVVATIKSVRFTEKNGQLIYPTFTDTPSVQRL 130
Qy 61 NSVLTLMISVIVVMTIFLVLYKYRCYKFIHGLWLMSSMLLFTYIYLG 113
Db 131 HSILNAINISVIVVMTIFLVLYKYRCYKFIHGLWLMSSMLLFTYIYLG 178
RESULT 7
Qy 019737 PRELIMINARY; PRT; 406 AA.
AC 019737; Q22692;
DT 01-NOV-1996 (TReMBLrel. 01, Created)
DT 01-MAY-1999 (TReMBLrel. 10, Last sequence update)
DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
DE F22E10.5 PROTEIN.
GN F22E10.5
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
OC Rhabditina; Rhabditidae; Rhabditidae; Peloderinae; Caenorhabditis.
RN [1]
RP SEQUENCE FROM N.A.
RA GARDNER A.;
RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
DR EMBL; Z67882; CAA91804.1; .
DR EMBL; Z50797; CAA91804.1; JOINED.
DR EMBL; Z50797; CAA90677.1; .
DR EMBL; Z67882; CAA90677.1; JOINED.
DR PROSITE; PS00379; CDP_ALCOHOL_P_TRANSF; 1.
SQ SEQUENCE 406 AA; 45628 MW; 07336492 CRC32;

Query Match 5.9%; Score 113.5; DB 5; Length 406;
Best Local Similarity 24.7%; Pred. No. 0.54;
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Db 136 HGCDSTQVFTLNICAYMSLGTVPVGLIVSVISVVMFYCAHWSYCTGQLRFSKFDVT 195
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Matches 40; Conservative 35; Mismatches 50; Indels 49; Gaps 8;

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QY      58 RLNSVLNTLIMISVIIVMTIFLVLYKYRCKYFIHGWLIMSSLMMLFLFYIYLGEVLK 117
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QY      118 TYNVANDYPTELLTVTNFGAVGMVCIIHWKGPLVLQQAYLIMISALMLVFTKYL 171
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Date: Mar 18, 2000 2:37 PM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-1998 CompuGen Ltd.

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Query length: 448

Database: GenEmbl.*

Database sequences: 821193

Database length: 1518192014

Search time (sec): 533.740000

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seq documentation block: 2285 bp DNA PAT 29-SEP-1999
LOCUS AR060156

DEFINITION Sequence 136 from patent US 5840540.

ACCESSION AR060156

VERSION AR060156.1 GI:5986606

KEYWORDS Unknown.

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 2285)

AUTHORS St. George-Hyslop,P.H., Rommens,J.M. and Fraser,P.E.

TITLE Nucleic acids encoding presenilin II

JOURNAL Patent: US 5840540-A 136 24-NOV-1998;

FEATURES Location/Qualifiers

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466 GCAGCAGGCGCCAGAGGATGAGAGATACTGCCAGTGGAGAGCCAG 515

51 GluAsnGluGluAspGlyGluGluAspProAspArgTyrValCysSerG1 67

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JOURNAL Submitted (16-DEC-1997) Molecular Biology, Tokyo Institute of
Psychiatry, 2-1-8 Kami-Kitazawa, Setagaya-ku, Tokyo 156, Japan
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AUTHORS	Sahara,N., Mori,H. and Shirasawa,T.				
TITLE	Molecular cloning of mouse presenilin 2 gene				
JOURNAL	Unpublished				
REFERENCE	2 (bases 1 to 1490)				
AUTHORS	Sahara,N., Mori,H. and Shirasawa,T.				
TITLE	Direct Submission				
JOURNAL	Submitted (16-DEC-1997) Molecular Biology, Tokyo Institute of Psychiatry, 2-1-8 Kami-kitazawa, Setagaya-ku, Tokyo 156, Japan				
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ACCESSION X99267

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REFERENCE Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
AUTHORS Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
JOURNAL 1 (bases 1 to 2088)
AUTHORS Frentzel, S., Abdel, A.S. and Luebbert, H.
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 2088)
AUTHORS Frentzel, S.
TITLE Direct Submission
JOURNAL Submitted (05-JUL-1996) S. Frentzel, Sandoz Pharma Ltd, Preclinical
RESEARCH, PO Box, CH-4002 Basel, Switzerland
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complete cds.

ACCESSION U34349

VERSION U34349.1 GI:1079575

KEYWORDS Alzheimer's disease.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominiidae; Homo.

REFERENCE 1 (bases 1 to 1673)

AUTHORS Li, J., Ma, J. and Potter, H.

TITLE Identification and expression analysis of a potential familial

Alzheimer disease gene on chromosome 1 related to AD3

Proc. Natl. Acad. Sci. U.S.A. 92 (26), 12180-12184 (1995)

JOURNAL 96109229

MEDLINE 2 (bases 1 to 1673)

AUTHORS Li, J.

TITLE Direct Submission

JOURNAL Submitted (21-AUG-1995) Jinhe Li, Neurobiology, Harvard Medical

School, 220 Longwood Ave., B2-502, Boston, MA 02115, USA

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AUTHORS TITLE JOURNAL

Tanahashi, H.
Direct Submission
Submitted (28-MAY-1997) to the DDBJ/EMBL/GenBank databases. Hiroshi
Tanahashi, National Institute of Neuroscience, Division of
Demyelinating Disease and Aging; 4-1-1 Ogawahigashi, Kodaira, Tokyo
187, Japan (E-mail: tanahashi@ncnp.nicp.go.jp, Tel: 81-423-41-1717,
Fax: 81-423-46-1747)

REFERENCE

2 (sites)

AUTHORS

Tanahashi, H. and Tabira, T.

JOURNAL

Biochim. Biophys. Acta 1396 (3), 259-262 (1998)

MEDLINE

98207716

COMMENT

Sequence updated (06-Jun-1997)

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VERSION A63557.1 GI:3717212
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SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 2681)
AUTHORS Talerman,A., Anson,R. and Cohen,D.
TITLE NUCLEOTIDE SEQUENCES, PROTEINS, DRUGS AND DIAGNOSTIC AGENTS FOR
TREATING CANCER
JOURNAL Patent: WO 9722695-A 10 26-JUN-1997;
COMMENT FONDATION JEAN DAUSSET CEPH (FR)
Other publication FR 2747691 19971024
Other publication FR 2742766 19970627.
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811 TTAAGACCTCAATGTCGCGTGACTACGTACAGTAGCAGCTCTCTAATC 860
209 TrpAsnPheGlyAlaValGlyMetValCysIleHisTrpLysGlyProLe 225
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
861 TGGAAATTTGGTGTGGTGGGATGATGCCATCCATCGAAGGCCCTCT 910
225 uValLeuGlnAlaTyrLeuIleMetIleSerAlaLeuMetAlaLeuV 242
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911 TCGACTCGAGCAGCGCTATCTCATTTATGATCAGTGCCTCATGGCCCTGG 960
242 alPheIleLysTyrLeuProGluTrpSerAlaTrpValIleLeuGlyAla 258
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
961 TATTATCAAGTACCTCCCGAATGGACCGCATGGCTCATCTTGGCTGTG 1010
259 IleSerValTyrAspLeuValAlaValLeuCysProLysGlyProLeuAr 275
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1011 ATTTCAGTATATGATTGGTGGCTGTTTATGTCCTCCAAAGGCCACTTCG 1060
275 gMetLeuValGluThrAlaGlnGluArgAsnGluProIlePheProAlaL 292
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1061 TATGCTGTTGAAACAGCTCAGGAAAGAAATGAGACTCTCTTTCCAGCTC 1110
292 euIleTyrSerSerAlaMetValTrpThrValGlyMetAlaLysLeuAsp 308
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1111 TTATCTATTCCTCAACAATGGTGTGGTGTGTAATATGCTGAAGGAGAC 1160
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|||||CCAGAACCCCAA.....AGGAGGGTACCCCAAGAACCCCAAGTATACAC 1135
325 .....GluAspSerTyrAspSerPheGlyGluP 334
1136 ACAAGAGCGGAGAGAGACACAGGACAGTGGTCTGGGAACGATGATG 1185
334 roSerTyrProGluValPheGluProProLeuThrGlyTyrProGly... 349
1186 GTGGCTTCAGTGAGAGTGGAGGCCCAAGAGACAGTCACTCGGGCGCT 1235
350 .....GluGluLeu..... 352
1236 CATCGCTCCACTCCCGAGTCAAGAGCTGCTGTCAGGAACCTTCTGGGAG 1285
353 .....GluGluGluGluArgGlyValIlystLeuGlyLeuG 365
1286 CATCTAACGAGTGAAGACCCGGAGGAAGAGGAGTAAACTTGGACTGG 1335
365 lyAspPheIlePheTyrSerValLeuValGlyLysAlaAlaAlaThrGly 381
1336 GAGATTTCATTTCTACAGTCTTCTGGTGGTAAAGGCTCAGCAACCCG 1385
382 SerGlyAspTrpAsnThrThrLeuAlaCysPheValAlaIleLeuIleG 398
1386 AGTGGAGACTGGAAACAACATAGCTGCTTTGTAGCATACTATCGG 1435
398 yLeuCysLeuThrLeuLeuLeuAlaValPheIlystLysAlaLeuProA 415
1436 CCTGTGCTTACATTACTCTGCTGCGCATTTTCAAGAAAGCGTGGCAG 1485
415 laLeuProIleSerIleThrPheGlyLeuIlePheTyrPheSerThrAsp 431
1486 CCTCCCATCTCCATCCTCGGGTCTGGTCTTACTTCGCCACGGAT 1535
432 AsnLeuValArgPropheMetAspThrLeuAlaSerHisGlnLeuTyrI 448
1536 TACCTGTGAGCGCTTCATGGACCAACTTGCATTCATCATGTTTATAT 1585
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1586 c 1586
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seq_name: gb_ro:MUSSI1PR

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seq_documentation_block: 1964 bp mRNA ROD 24-JUL-1995
LOCUS MUSSI1PR Mus musculus S182 protein mRNA, complete cds.
DEFINITION ACCESSION L42177
VERSION L42177.1 GI:904129
KEYWORDS S182 protein.
SOURCE Mus musculus
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 1964)
AUTHORS Sherrington,R., Rogava,E.I., Liang,Y., Rogava,E.A., Levesque,G.,
Ikeda,M., Chi,H., Lin,C., Li,G., Holman,K., Tsuda,T., Mar,L.,
Foncin,J.-F., Bruni,A.C., Montesi,M.P., Sorbi,S., Rainero,I.,
Pinassi,L., Nee,L., Chumakov,I., Pollen,D., Brookes,A., Sanseau,P.,
Polinsky,R.J., Masco,W., Da Silva,H.A.R., Haines,J.L.,
Perlick-Vance,M.A., Tanzi,R.E., Roses,A.D., Fraser,P.E.,
Romans,J.M. and St. George-Hyslop,P.H.
Cloning of a gene bearing missense mutations in early-onset
familial Alzheimer's disease
NATURE 375 (6534), 754-760 (1995)
JOURNAL 95319502
MEDLINE
COMMENT On Jul 25, 1995 this sequence version replaced gi:897616.
FEATURES Location/Qualifiers
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source
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/db_xref="taxon:10090"
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/map="14q24.3"
/note="clone: composite sequence from a ~ 0.7 kb RT-PCR
clone and an overlapping ~2.0 kb cDNA clone mc6-3-2
recovered from adult mouse brain cDNA library in
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5'UTR
mRNA
CDS
1..187
1..1964
188..1591
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number L42110); putative"
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/product="S182 protein"
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VIHAWLLISLLLLFFESFYILSEVKTYNVAVDYTVALLINFGVGMIAIHWKGP
LRQAYLIMISALMALVFIKLPETAWLILAVISYDLVAVLCPPKPLRMLVETAQ
ERNETLFPALYISSTMVNLVMAEGDPEAQRVRPKYNTQRAERTQDSGSGNDG
GFSEWEAQDRLSHLPHSTPESRAAVQELSGILTSDEPERGVKLGDFIFYSVL
VGKASATAGMDNTIACFVAILIGLCLTLLLAIFKALPALPISITFGLVVFATD
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BASE COUNT 503 a 503 c 496 g 460 t 2 others
ORIGIN
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alignment_scores:
Quality: 1468.00 Length: 467
Ratio: 3.915 Gaps: 6
Percent Similarity: 80.300 Percent identity: 64.026
alignment_block:
US-08-509-359B-137 x MUSSI1PR
Align seg 1/1 to: MUSSI1PR from: 1 to: 1964
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24 GluSerProThrProArgSerCysGlnGluGlyArgGlnGlyProGluAs 40
194 GAGATACCTGCACCTTTCTCTACTTCCAGATGCCAGATGCTGTGAGGA 243
40 pGlyGluAsnThrAlaGlnTrpArgSerGlnGluAsnGluGluAspGlyG 57
244 CAGCCACTCCAGCAGCGCCATCCGGAGCCAGATGACAGCAAGACGCGC 293
57 luGluAsProAspArgTyrValCysSerGlyValPro..... 69
294 AGCAGCAGCATGACGAGCAGACTTGACACCCCTGAGCCAATATCTAAT 343
70 GlyArgProProGly.....Le 75
344 GGGCGGCCCCAGAGTAACCAAGACAGGTGGTGGAAACAGATGAGGAGGA 393
75 uGluGluLeuThrLeuLysTyrGlyAlaLysHisValIleMetLeuP 92
394 AGACCAAGAGCTGACATTAATATGAGCAAGCATGTCATCATGCTCT 443
92 heValProValThrLeuCysMetIleValValAlaThrIleLysSer 108
444 TTGTCCCGTGCACCTCTGTCATGTCGTCGTCGTCGTCGTCGTCGTCGTC 493
109 ValArgPheTyrThrGluLysAsnGlyGlnLeuIleTyrThrProPheTh 125
494 GTCAGCTTCTATACCCGGAAGACGGTCAGCTAACTACACCCCATTCAC 543
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CC acid probes specific for the mutant gene, provides a means of
 CC diagnosing Alzheimer's disease.
 SQ Sequence 2236 BP; 488 A; 584 C; 645 G; 519 T;

alignment_scores:

Quality: 2336.00 Length: 448
 Ratio: 5.214 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-08-509-359b-137 x T51253

Align seg 1/1 to: T51253 from: 1 to: 2236

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1 MetLeuThrPheMetAlaSerAspSerGluGluValCysAspGluAr 17
368 ATGCTCACATTCATGGCCCTCTGACAGCGAGGAAGTGTGTATGAGCG 417
17 gThrSerLeuMetSerAlaGluSerProThrProArgSerCysGlnGlu 34
418 GAGTCCCTTAATGTGGCGGAGAGCCCGCCGCGCTCTCGCCAGGAGG 467
34 lyArgGlnGlyProGluAspGlyGluAsnThrAlaGlnTrpArgSerGln 50
468 GCAGGCAGGGCCAGAGGATGGAGAACACATGCCCCAGTGGAGAGCCAG 517
51 GluAsnGluGluAspGlyGluGluAspProAspArgTyrValCysSerGln 67
518 GAGAACGAGGAGGCGGTGAGGAGGACCTGACCGCTATGCTGTAGTGG 567
67 yValProGlyArgProProGlyLeuGluGluLeuThrLeuLysTyrG 84
568 GGTTCCTCCGGCGCGCCCGAGGCTGGAGGAAGAGCTGACCCCTCAATACG 617
84 lyAlaLysHisValIleMetLeuPheValProValThrLeuCysMetIle 100
618 GAGCGAAGCACGTGATCATGCTGTTTGTGCGTGTCACTCTGTGTCATGATC 667
101 ValValValAlaThrIleLysSerValArgPheTyrThrGluLysAsnGln 117
668 GTGGTGGTAGCCACCATCAAGTGTGTGCGCTTCTACACAGAGAAGATGG 717
117 yGlnLeuIleTyrThrProPheThrGluAspThrProSerValGlyGluA 134
718 ACAGCTCATCTACAGCCCATCTACTGAGGACACACCCCTCGGTGGGCGAGC 767
134 rgLeuLeuAsnSerValLeuAsnThrLeuIleMetIleSerValIleVal 150
768 GCCTCTCACTCCGTCGTCGACACCCCTCATCATGATGATGCGTCATCGTG 817
151 ValMetThrIlePheLeuValValLeuTyrLysTyrArgCysTyrLysPh 167
818 GTTATGACCATCTTCTGTGTGTGTCCTACAAGTACCGCTGCTACAAGTT 867
167 eileHisGlyTrpLeuIleMetSerSerLeuMetLeuLeuPheLeuPheT 184
868 CATCCATGGCTGGTGTGATCATGCTCTTCACTGATGCTGCTTCTCTTCA 917
184 hrTyrIleTyrLeuGlyGluValLeuLysThrTyrAsnValAlaMetAsp 200
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201 TyrProThrLeuLeuLeuThrValTrpAsnPheGlyAlaValGlyMetVa 217
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217 lCysIleHisTrpLysGlyProLeuValLeuGlnGlnAlaTyrLeuIleM 234
1018 GTGCATCCACTGGAAGGGCCCTCTGTGTGTGTCAGCAGGCGCTACCTCATCA 1067
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1068 TGATCAGTGGCGCTCATGGCCCTAGTGTTCATCAAGTACCTCCACAGATGG 1117
251 SerAlaTrpValIleLeuGlyAlaIleSerValTrpAspLeuValAlaVa 267
1118 TCCGCGTGGGTCTATCTCTGGCGCCATCTCTGTGTATGATCTCGTGGCTGT 1167
267 lLeuCysProLysGlyProLeuArgMetLeuValGluThrAlaGlnGluA 284
1168 GCTGTGTCCTCCAAAGGGCCTCTGAGAATGCTGGTAGAAGTCTCCACAGGAGA 1217
284 rgAsnGluProIlePheProAlaLeuIleTyrSerSerAlaMetValTrp 300
1218 GAAATGAGCCCATATTCCTGCGCTGATATATCTCATCTGCCATGTGTGG 1267
301 ThrValGlyMetAlaLysLeuAspProSerSerGlnGlyAlaLeuGlnLe 317
1268 ACGGTTGGCATGGCAAGCTGGACCCCTCTCTCAGGGTGGCCCTCCAGCT 1317
317 uProTyrAspProGluMetGluGluAspSerTyrAspSerPheGlyGluP 334
1318 CCCTAGCACCCGAGATGGAAGAGACTCTCTATGACAGTTTGGGGAGC 1367
334 roSerTyrProGluValPheGluProProLeuThrGlyTyrProGlyGlu 350
1368 CTTCATACCCCGAAGTCTTTGAGGCTCCCTTGACTGGCTACCCAGGGAG 1417
351 GluLeuGluGluGluGluArgGlyValLysLeuGlyLeuGlyAspPh 367
1418 GAGCTGGAGGAGGAGGAGGAGGCGGTGAAGCTTGGCCTCGGGGACTT 1467
367 eilePheTyrSerValLeuValGlyLysAlaAlaIleLeuIleGlyLeu 384
1468 CATCTTCTACAGTGTCTGGTGGCAAGGCGGTGCCACGGGCGAGCGGG 1517
384 spTrpAsnThrThrLeuAlaCysPheValAlaIleLeuIleGlyLeu 400
1518 ACTGGAATACACGCTGGCTGCTCTGCTGGCCATCCTCATTTGGCTGTGT 1567
401 LeuThrLeuLeuLeuLeuAlaValPheLysLysAlaLeuProAlaLeuPr 417
1568 CTGACCCCTCTGCTGCTGTGTGTGTTCAAGAAAGCGCTGCCCGCCCTCCC 1617
417 oileSerIleThrPheGlyLeuIlePheTyrPheSerThrAspAsnLeu 434
1618 CATCTCCATCACGTTCCGGGCTCATCTTTTACTTCTCCACGGACACCTGG 1667
434 alArgProPheMetAspThrLeuAlaSerHisGlnLeuTyrIle 448
1668 TGGCGGCGTTCATGGACACCCCTGGCCCTCCCATCAGCTCTACATC 1711

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seq_name: N_Geneseq_36:V04669

seq_documentation_block:

ID V04669 standard; CDNA; 2229 BP.
 AC V04669;
 DT 20-JUL-1998 (first entry)
 DE Human presenilin-2 cDNA (hps2).
 KW Presenilin-1; PS1 gene; human; familial Alzheimer's disease; FAD;
 KW cerebral haemorrhage; schizophrenia; depression; epilepsy;
 KW mental retardation; diagnosis; therapy; transgenic animal; ss.
 OS Homo sapiens.
 FH Key
 Location/Qualifiers
 366..1715
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 /tag= p
 /note= "A to T FAD mutation site (Asn141Ile)"
 mutation 1080
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 mutation 1624
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1616 CATCTCCATCACGCTCGGCTCATCTTTTACTTCTCCACGAGCAACCTGG 1665
 434 aArgProPheMetAspThrLeuAlaSerHisGlnLeuTyrIle 448
 1666 TGGCGCGCTTCATGGACACCTGGCCCTCCCATCAGCTCTACATC 1709

seq_name: N_Geneseq_36.T87426

seq_documentation_block:

ID T87426 standard; DNA; 2276 BP.

AC T87426;

DT 07-DEC-1997 (first entry)

DE Full AD4/AD3LP sequence.

KW AD3: AD4/AD3LP: Alzheimer's disease; chromosome; missegregation;

KW presenilin; inhibitor; AD; trisomy 21; ss.

OS Homo sapiens.

PN WO9707213-A2.

PD 27-FEB-1997.

PF 15-AUG-1996; U13114.

PR 16-AUG-1995; US-002448.

PA (HARD) HARVARD COLLEGE.

PI Li J., Potter H;

DR WPI; 97-165297/15.

DR P-PSDB; W28508.

PT Identifying genes which cause chromosome missegregation - useful for

PT identifying causes of and treatments for diseases, e.g. Alzheimer's

PS Claim 28; Fig 28; 7pp; English.

CC Identifying genes which cause improper chromosome segregation,

CC screening for inhibitors of chromosome missegregation and processes

CC caused by genes encoding chromosome missegregation promoters

CC was exemplified using Alzheimer's disease. The sequences

CC given in T87401 to T87426 can be used in the above methods.

CC It is not clear from the figure legend, the figure and the

CC disclosure of the specification which sequence of Fig 1 and Fig 28

CC is the AD4/AD3LP or the AD3 sequence.

SQ Sequence 2276 BP; 494 A; 595 C; 562 G; 525 T;

alignment_scores:

Quality: 2320.50

Ratio: 5.191

Percent Similarity: 99.777

Percent Identity: 99.777

alignment_block:

us-08-509-359B-137 x T87426 ..

Align seg 1/1 to: T87426 from: 1 to: 2276

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 461 GAGCTCCCTAATGTGGCCGAGAGCCCCACGCGCGCTCTGCGAGGAGG 510
 34 lyArgGlnGlyProGluAspGlyGluAsnThrAlaGlnTrpArgSerGln 50
 511 GCAGGACGGGCCAGAGATGGAGAGACACTGCCAGTGGAGAGCCAG 560
 51 GluAsnGluGluAspGlyGluGluAspProAspArgTyrValCysSerGl 67
 561 GAGAACGAGGAGCGGTGAGGAGGACCTGACCGCTATGCTCTAGTGG 610
 67 yValProGlyArgProGlyLeuGluGluGluLeuThrLeuLysTyrG 84
 611 GGTTCGCGCGCGCCGAGCGCTGGAGGAGAGCTGACCCCTCAAATACG 660
 84 lyAlaLysHisValIleMetLeuPheValProValThrLeuCysMetIle 100
 661 GACGAGACGAGGTGATGCTGTTGTGCTGTCACCTCTGTGTCATGATC 710

101 ValValValAlaThrIleLysSerValArgPheTyrThrGluLysAsnGl 117
 711 GTGGTGTAGCCACCATCAAGTCTGTGGCTTCTACACAGAGAAATGG 760
 117 yGlnLeuIleTyrThrProPheThrGluAspThrProSerValGlyGlnA 134
 761 ACAGCTCATCTACAGCCATTCACCTGAGGACACACCCCTCGGTGGCCAGC 810
 134 rgLeuLeuAsnSerValLeuAsnThrLeuIleMetIleSerValIleVal 150
 811 GCCTCCTCAACTCGGTGCTGAACACCCCTCATCATGATCATGAGGTATCGTG 860
 151 ValMetThrIlePheLeuValValLeuTyrLysTyrArgCysTyrLysPh 167
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 167 eileHisGlyTriPLeuIleMetSerSerLeuMetLeuLeuPheLeuPhe 184
 911 CATCATGGCTGGTGTGATCATGCTTCTCAGTATGCTGCTGTCTCTCTCA 960
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 251 SerAlaTrpValIleLeuGlyAlaIleSerValTyrAspLeuValAlaVa 267
 1161 TCCGCGTGGGTTCATCCTGGGCGCATCTCTGTATGATCTCGTGGCTGT 1210
 267 lLeuCysProLysGlyProLeuArgMetLeuValGluThrAlaGlnGluA 284
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 1261 GAATGAGCCCATATTCCTGCTCCCTGATATACTCATCTGCCATGGTGTGG 1310
 301 ThrValGlyMetAlaLysLeuAspProSerSerGlnGlyAlaLeuGlnLe 317
 1311 ACGTTGGCATGGCAAGCTGGACCCCTCTCTCAGGGTGGCCCTCCAGCT 1360
 317 uProTyrAspProGluMetGluAspSerTyrAspSerPheGlyGluP 334
 1361 CCCCTACGACCCCGAGATG...GAAGACTCCTATGACAGTCTTGGGGAGC 1407
 334 roSerTyrProGluValPheGluProProLeuThrGlyTyrProGlyGlu 350
 1408 CTTTCATCCCGAAGTCTTTGAGGCTCCCTTGACTGCTACCCAGGGGAG 1457
 351 GluLeuGluGluGluGluGluArgGlyValLysLeuGlyLeuGlyAspPh 367
 1458 GAGCTGGAGGAAGAGAGGAAAGGGCGTGAAGCTTGGCCCTCGGGGACTT 1507
 367 eilePheTyrSerValLeuValGlyLysAlaAlaAlaThrGlySerGlyA 384
 1508 CATCTTCTACAGTGTGTTGGTGGCAAGCGGCTGCCACGCGGACGGGG 1557
 384 spTrpAsnThrThrLeuAlaCysPheValAlaIleLeuIleGlyLeuCys 400
 1558 ACTGGAATACCACCTGGCCCTGCTTGTGGCCATCTCTATTGGCTTGTGT 1607

seq_name: N_Geneseq_36:T40030

seq_documentation_block:

ID T40030 standard; DNA; 1964 BP.

AC T40030:

DT 23-JUL-1997 (first entry)

DE Murine presenilin-1 wild type coding sequence.

KW Presenilin-1; mouse; hpsl-1; hpsl-2; PS-2; integral membrane protein; AD;

KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;

OS depression; antibody; gene expression modulator; therapy; ss.

FT Key Location/Qualifiers

FT 188..1591

FT cds

FT /*tag= a

FT /product= presenilin-1

PN W09634099-A2.

PD 31-OCT-1996.

PF 29-APR-1996; CA0263.

PR 28-APR-1995; US-431048.

PR 28-JUN-1995; US-496841.

PR 31-JUL-1995; US-509359.

PA (HSCR-) HSC RES & DEV LP.

PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.

PI Fraser PE, Rommens JM, St George-Hyslop PH;

DR WPI; 96-497631/49.

DR P-PSDB; W05735.

PT New presenilin genes - useful for diagnosis, therapy and drug

PT screening of familial Alzheimer's disease, cerebral disorders, etc.

PS Claim 8; Page 145-146; 178pp; English.

CC This sequence represents the coding sequence for the murine presenilin-1.

CC T40028 and T40029 represent the coding sequences for the two different

CC forms of wild type human presenilin-1 (PS-1). The form represented by

CC T40029 results from alternate splicing of the genomic DNA sequence.

CC T40031 represents the coding sequence for wild type human PS-2. The

CC presenilins are a family of highly conserved integral membrane proteins

CC with a common structural motif, common alternate splicing patterns, and

CC common mutational hot spot regions. Mutations in PS genes are implicated

CC in familial Alzheimer's disease (AD) and possibly other diseases such as

CC cerebral haemorrhage, schizophrenia, depression etc., so detection of

CC mutations in these sequences can be used for diagnosis of these diseases.

CC The encoded proteins, or vectors that express them or containing

CC antisense sequences, antibodies selective for mutant forms of the encoded

CC proteins (such as W05736) and modulators of PS gene expression are useful

CC as models for drug screening. The antibodies can also be used e.g. for

CC affinity purification and in immunoassays.

SQ Sequence 1964 BP; 503 A; 503 C; 496 G; 460 T;

alignment_scores:

Quality: 1468.00 Length: 467

Ratio: 3.915 Gaps: 6

Percent Similarity: 80.300 Percent Identity: 64.026

alignment_block:

US-08-509-359b-137 x T40030 ..

Align seg 1/1 to: T40030 from: 1 to: 1964

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194 GAGATACCTCGACCTTTGCTTCTACTTCCAGAAATGCCAGATGCTCTGAGGA 243

40 pGlyGluAsnThrAlaGlnTrpArgSerGlnGluAsnGluGluAspGlyG 57

244 CAGCCACTCCAGAGCGCCATCGGAGCCAGATGACAGCCAGAACGGC 293

57 luGluAspProArgTyrValCysSerGlyValPro..... 69

294 AGCAGCAGCATGACAGCGCAGACTTGACACACCTGAGCCAATATCTAAT 343

70 GlyArgProProGly.....Le 75

344 GGGGGCCCCCAGAGTAACACTCAAGACAGGTGGTGAACAAGATGAGGAGGA 393

75 uGluGluGluLeuThrLeuLysTyrGlyAlaLysHisValIleMetLeuP 92

394 AGCGAAGAGCGTGACATGAAATATGGAGCCCAAGCATGTCATCATGCTCT 443

92 heValProValThrLeuCysMetIleValValValAlaThrIleLysSer 108

444 TTGTCGCCCGTGACCTCTGCAATGTCGTCGTCGTCGCCACCATCAATCA 493

109 ValArgPheTyrThrGluLysAsnGlyGlnLeuIleTyrThrProPheTh 125

494 GTCAGCTTCTATACCCGGAAGGCGGTACAGCTAATCTACACCCCATTCAC 543

125 rGluAspThrProSerValGlyGlnArgLeuLeuAsnSerValLeuAsnT 142

544 AGAAGACACTGAGACTGTAGCCCAAGAGCCCTGCACCTCGATCCTGAATG 593

142 hrLeuIleMetIleSerValIleValValMetThrIlePheLeuValVal 158

594 CGGCATCATGATCAGTGTCAATTGTATTATGACCATCCTCTGGTGGTC 643

159 LeuTyrLysTyrArgCysTyrLysPheIleHisGlyTrpLeuIleMetSe 175

644 CTGTATAAATACAGTGCTACAAGGTCAATCCAGCGCTGCGTATTATTTC 693

175 rSerLeuMetLeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluValL 192

694 ATCTCTGTGTGTGTTCTTTTTCGTTTCTATTACTTAGGGGAAGTAT 743

192 euLysThrTyrAsnValAlaMetAspTyrProThrLeuLeuThrVal 208

744 TTAAGACTCAAAATGTCGCGTGGATACGTACGTAGTACACTCTCAATC 793

209 TrpAsnPheGlyAlaValGlyMetValCysIleHisTrpLysGlyProLe 225

794 TGGAAATTTGGTGGTGGGATGATTCCTCCATCCACTGGAAGGCCCTCT 843

225 uValLeuGlnAlaTyrLeuIleMetIleSerAlaLeuMetAlaLeuV 242

844 TCGACTGCAGCAGCGGTATCTCATTATGATCAGTCCCTCATGGCCCTGG 893

242 alPheIleLysTyrLeuProGluTrpSerAlaTrpValIleLeuGlyAla 258

894 TATTATCAAGTACTCTCCCAATGGACCGCATGCTCATCTTGGCTGTG 943

259 IleSerValTyrAspLeuValAlaValLeuCysProLysGlyProLeuAr 275

944 ATTTCAGTATATGATTTGGTGGCTGTTTATGTCCTCCAAAGGCCACTTCG 993

275 gMetLeuValGluThrAlaGlnGluArgAsnGluProIlePheProAlaL 292

994 TATGCTGTTTGAACAGCTCAGGAAGAAATGAGACTCTCTTCCAGCTC 1043

292 euIleTyrSerSerAlaMetValTrpThrValGlyMetAlaLysLeuAsp 308

1044 TTATCTATTCCTCAACAATGTTGTTGGTGAATATGCTGAAGGAGAC 1093

309 ProSerSerGlnGlyAlaLeuGlnLeuProTyrAspProGluMetGlu... 324

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325GluAspSerTyrAspSerPheGlyGluP 334

1138 ACAAGACCGGAGAGACAGACAGGACAGTGGTCTCGGAACGATGATG 1187

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1238 CATCGCTCCACTCCCGAGTCAAGAGCTGCTGTCCAGGAACCTTCTGGGAG 1287


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91 euPheValProValThrLeuCysMetileValValValAlaThrileLys 107
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158 ValLeuTyrLysTyrArgCysTyrLysPheIleHisGlyTrpLeuIleMe 174
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174 tSerSerLeuMetLeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluV 191
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952 TGGTGTATTATCAAGTACCTCCCTGATGAGTGCCTGGCTCATCTTGGCT 1001
258 AlaIleSerValTyrAspLeuValAlaValLeuCysProLysGlyProLe 274
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1002 GTGATTTTCAGTATGATTTAGTGGCTGTTTGTGTCGGAAGGTCCACT 1051
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308 AspProSerSerGlnGlyAlaLeu.....GlnLeuProTyrAspProGl 322
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ID V17358 standard; DNA; 2764 BP.
AC V17358;
DT 04-JUN-1998 (first entry)
DE PS1/467 protein coding sequence.
KW Presenilin peptide; PS1/429; immunogen; immune response; PS1 gene;
KW Alzheimer's disease; mitochondrial pathology; neurodegeneration;
KW apoptosis; PS1/467; ss.
OS Homo sapiens.
FH Key Location/Qualifiers
FT CDS 249..1652
   /tag= a
   WO9746678-Al.
   PD 11-DEC-1997.
   PF 03-JUN-1997; U09272.
   PR 18-JUL-1996; US-683315.
   PR 06-JUN-1996; US-659296.
   PA (FARB ) BAYER CORP.
   PI Chisholm JC, Davis JN, Drache B;
   DR WPI: 98-042186/04.
   DR P-PSDB; W41430.
   PT DNA encoding presenilin peptide PS1/429 and its analogues - useful
   for diagnosis and treatment of Alzheimer's disease
   PS Disclosure; Fig 2; 77pp; English.
   CC This sequence encodes the PS1/467 presenilin peptide. This sequence is
   specifically stated as not being in the nucleic acid of the invention,
   which encodes the PS1/429 presenilin peptide PS1/429 (ii). Cells
   transformed with the DNA are used to produce recombinant (ii) and
   analogues, useful e.g. as immunogens for generating an immune response
   against PS1/429. (ii) is a new product of the PS1 gene, mutations in
   which cause Alzheimer's disease (AD). The nucleic acids are generally
   useful as probes for detection and quantification of PS1/429,
   particularly for diagnosis of AD, especially the target sequences that
   hybridize with probes are isolated for sequencing. Antibodies (Ab) can
   also be diagnosed at the protein level using Ab as immunoassay reagents.
   Ab can also be used to identify epitopes and for affinity purification of
   peptides. Antisense nucleic acid may also be used to regulate expression
   of the PS1/429 gene, and both nucleic acids and peptides are useful as
   size markers in electrophoresis, chromatography etc. The transgenic
   animals are used as models for AD, e.g. for testing drugs. Regulators of
   the PS1/429 gene or polypeptide can be used to treat e.g. AD or diseases
   involving mitochondrial pathology, apoptosis and neurodegeneration.
   CC Typical regulators are antisense sequences, ribozymes, aptamers,
   synthetic or natural compounds. (ii) may also be used to target other
   coding sequences to particular cellular locations.
   CC Sequence 2764 BP; 715 A; 624 C; 653 G; 772 T;
   SQ

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[illegible]

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40  pGly.....GluAsnThrAlaGlnTrpArgSerGlnGluAsnGluGluA 55
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230  CAACCACTGCAGCAATCTNNNNNN...NNNNNNATGACAATAGAGAC 276
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277  GGCAGAGCAGCAACAGCAGCGGCTTTGGCCACCCTGAGCCATTATCT 326
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70  ...GlyArgProGly..... 74
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327  AATGGAGCCCCAGGGTAACCTCCCGCAGGTGGTGACCAAGATGAGGA 376

75  LeuGluGluGluLeuThrLeuLysTyrGlyAlaLysHisValIleMetL 91
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377  AGAAGATGAGGAGCTCACATTGAAATATGGCCCAAGCATGTGATCATCC 426

91  eupheValProValThrLeuCysMetIleValValAlaThrIleLys 107
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427  TCTTTTGCCGTGTGACTCTGCATGGTGGTGGTGGTGGTGGTGGTGGT 476
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350 .....GluGluLeu..... 352
1221 ACACCTGAGTCACGAGCTGTCTCCAGGAACCTTCCACAGTATCCTCGC 1270
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353 .....GluGluGluGluGluArgGlyValLysLeuGlyLeuGlyAspPheI 368
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368 lePheTyrSerValLeuValGlyLysAlaAlaAlaThrGlySerGlyAsp 384
    ||| :|||
1321 TTTTCTACAGTGTCTGTTGGTTAAAGCCTCAGCACAGCCAGTGGAGAC 1370
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385 TrpAsnThrThrLeuAlaCysPheValAlaAlaLeuLeuGlyLeuCysLe 401
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1371 TGGAACACACCATAGCCTGTTTCGTAGCCATATTAATTGTTTGCCT 1420
    ||| :|||
401 uThrLeuLeuLeuAlaValPheLysLysAlaLeuProAlaLeuProI 418
    ||| :|||
1421 TACATTATTACTCTTCCATTTTCAAGAAAGCATTCGCCAGCTCTCCAA 1470
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418 leSerilleThrPheGlyLeuIlePheTyrPheSerThrAspAsnLeuVal 434
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1471 TCTCCATCACCTTTGGCTTGTCTTCTACTTTGCCACAGATTATCTTGT 1520
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Align seg 1/1 to: US-08-592-541-136 from: 1 to: 2285

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416 GACGTCCCTAATGTTCGGCCGAGAGCCCCACGCCGCTCTCCAGGAGG 465
34 lyArgGlnGlyProGluAspGlyGluAsnThrAlaGlnTrpArgSerGln 50
466 GCAGGCGAGGCCCCAGAGGATGTGAGAGAATACTGCCAGTGGAGAAGCCAG 515
51 GluAsnGluGluAspGlyGluGluAspProAspArgTyrValCysSerGln 67
516 GAGAACGAGGAGGAGCGGTGAGGAGGACCTGACCGCTATGTCTGTAGTGG 565
67 yValProGlyArgProProGlyLeuGluGluLeuThrLeuLysTyrG 84
566 GTTCCCGGGGGCCGCGAGGCTGGAGGAGAGCTGACCCCTCAAAATACG 615
84 lyAlaLysHisValIleMetLeuPheValProValThrLeuCysMetIle 100
616 GAGCGAAGCATGTGATCATCTGTTTGTGCTGTCACTCTGTGATGATC 665
101 ValValValAlaThrIleLysSerValArgPheTyrThrGluLysAsnGln 117
666 GTGGTGTAGCCACCATCAAGTCTGTGGCTTCTACACAGAGAAGATGG 715
117 yGlnLeuIleTyrThrProPheThrGluAspThrProSerValGlyGlnA 134
716 ACAGCTCATCTACAGCGCATCTACGTGAGGACACACCCCTCGGTGGGCCAGC 765
134 rgLeuLeuAsnSerValLeuAsnThrLeuIleMetIleSerValIleVal 150
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151 ValMetThrIlePheLeuValValLeuTyrLysTyrArgCysTyrLysPh 167
816 GTTATGACCATCTTCTTGTGTGTGCTCTACAAGTACCGCTGCTCAAGTT 865
167 eIleHisGlyTrpLeuIleMetSerSerLeuMetLeuLeuPheLeuPheT 184
866 CATCCATGGGTGTGTGATCATGCTTCACTGATGCTGCTGCTCTCTCA 915
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916 CCTATATCTACCTTGGGAGTGTCTGGAACCTCAAGACCTAAGTGGCCATGGAC 965
201 TyrProThrLeuLeuLeuThrValTrpAsnPheGlyAlaValGlyMetVa 217
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1016 GTGCATCCATCGGAAGGCGCTCTGGTGTCTGACGAGCGCTACCTCATCA 1065
234 eIleSerAlaLeuMetAlaLeuValPheIleLysTyrLeuProGluTrp 250
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251 SerAlaTrpValIleLeuGlyAlaIleSerValTyrAspLeuValAlaVa 267
1116 TCCGCGTGGGTATCATCTGGGGCCATCTCTGTGTATGATCTCGTGGCTGT 1165
267 lLeuCysProLysGlyProLeuArgMetLeuValGluThrAlaGlnGluA 284
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334 roSerTyrProGluValPheGluProProLeuThrGlyTyrProGlyGlu 350
1366 CTTATACCCCGAAGCTTTGAGCTCCCTTACTGGCTACCCAGGGAG 1415
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1416 GAGCTGGAGGAGGAGGAGGAGGCGCTGAAGCTTGGCTCGGGACTT 1465
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; Sequence 28, Application US/08875972
; Patent No. 5985564
; GENERAL INFORMATION:
; APPLICANT: Huntington Potter and Jinhue Li
; TITLE OF INVENTION: ASSAY FOR IDENTIFYING GENES CAUSING
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HAMILTON, BROOK, SMITH & REYNOLDS, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02173-4799
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/875,972
; FILING DATE: 08-AUG-97
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/002,448
; FILING DATE: 16-AUG-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan Esq., Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: HU95-03PA
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781) 861-6240
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; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/875,972
; FILING DATE: 08-AUG-97
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/002,448
; FILING DATE: 16-AUG-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan Esq., Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: HU95-03PA
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781) 861-6240
; TELEFAX: (781) 861-9540
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1417 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: Double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 2..1129
; US-08-875-972-1

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; Sequence 135, Application US/08967101
; Patent No. 5840540
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street tower - 125 High Street
; CITY: Boston

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; Sequence 135, Application US/08592541
; Patent No. 5986054
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSES: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/592,541
; FILING DATE:
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 135:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1962 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-592-541-135

alignment_scores:
Quality: 1468.00 Length: 467
Ratio: 3.915 Gaps: 6
Percent Similarity: 80.300 Percent Identity: 64.026

alignment_block:

US-08-509-359B-137 x US-08-592-541-135

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192 GAGATACCTGCACCTTTGCTCTACTTCCAGATGCCAGATGCTGAGGA 241
40 pGlyGluAsnThrAlaGlnTrpArgSerGlnGluAsnGluGlyAspGly 57
242 CAGCCACTCCAGCAGCCCATCCGAGCCAGATGACAGCAAGACGGC 291
57 luGluAspProAspArgTyrValCysSerGlyValPro..... 69

292 AGCAGCAGCATGACAGGCAGAGACTTGACAACCTTGAGCCAATATCTAAT 341
70 GlyArgProProGly.....Le 75
342 GGGCGGCCCCAGAGTAACCTCAAGCAGGTGCTGGAACAAGATGAGGAGGA 391
75 uGluGluGluLeuThrLeuLysTyrGlyAlaLysHisValIleMetLeu 92
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92 heValProValThrLeuCysMetIleValValValAlaAlaThrIleLysSer 108
442 TTGTCCCGCTGACCTCTGCATGTGTCGTCGTGGCCACCATCAATCA 491
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492 GTCAGCTTCTATACCGGAGGAGCGGTACGCTAATCTACACCCATTCCAC 541
125 rGluAspThrProSerValGlyGlnArgLeuLeuAsnSerValLeuAsn 142
542 AGAAGACACTGAGACTGTAGGCCAAAGAGCCCTGCATCGATCTGAATG 591
142 hrLeuIleMetIleSerValIleValValMetThrIlePheLeuValVal 158
592 CGGCCATCATGATCAGTGTCTATTGTCATTATGACCATCTCTCTGTGTC 641
159 LeuTyrLysTyrArgCysTyrLysPheIleHisGlyTrpLeuIleMetse 175
642 CTGTATAAATACAGGTGCTACAAGGTCATCCACGCTGGCTTATTATTC 691
175 rSerLeuMetLeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluVal 192
692 ATCTGTGTGTGCTGCTTTTTCGTTTCTTACTTAGGGGAAGTAT 741
192 euLysThrTyrAsnValAlaMetAspTyrProThrLeuLeuThrVal 208
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209 TrpAsnPheGlyAlaValGlyMetValCysIleHisTrpLysGlyProle 225
792 TGAATTTTGGTGTGGTGGATGATGCCATCCACTGGAAGGCCCT 841
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842 TCGACTGCAGCAGCGGTATCTCATATGATCAGTGCCTCATGGCCCTGG 891
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259 IleSerValTyrAspLeuValAlaValLeuCysProLysGlyProLeuAr 275
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; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC Compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/592,541
; FILING DATE:
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 133:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2791 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: CDNA
; US-08-592-541-133

alignment_scores:
    Quality: 1467.00      Length: 466
    Ratio: 3.954          Gaps: 8
    Percent Similarity: 79.614 Percent Identity: 65.665

alignment_block:
US-08-509-359B-137 x US-08-592-541-133 ..

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305  CAACACCCCTGCAGCAATPACTGTATGACGTACCCAGAAATGACAAATGAGA 354

57  luGluAspProAspArgTyValCysSerGlyValPro..... 69
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355  AGGACACACACACACAGCG...AGCCTTGCCACCCCTGAGCCATTATCT 401

70  ..GlyArgProProGly..... 74
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402  AATGGACGACCCCGAGGTAACCTCCCGCAGGTGCTGGAGCAAGATGAGGA 451

75  .LeuGluGluGluLeuThrLeuLysTyrGlyAlaLysHisValIleMetL 91
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91  ePheValProValThrLeuCysMetIleValValAlaThrIleLys 107
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502  TCTTTGTCCTGTGACTCTCTCATGTGTTGCTGCTGCTGACCAATTAG 551

108  SerValArgPheTyrThrGluLysAsnGlyGlnLeuIleTyrThrProPh 124
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552  TCAGTCAGCTTTTATACCCGGAGGATGGCAGCTAATCTATACCCCAT 601

124  eThrGluAspThrProSerValGlyGlnArgLeuLeuAsnSerValLeuA 141
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141  snThrLeuIleMetIleSerValIleValValMetThrIlePheLeuVal 157
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652  ATGCTGTCCCATCATGATCAGTGTCTATTTGTCATGACTATCTCTCTGTTG 701

158  ValLeuTyrLysTyrArgCysTyrLysPheIleHisGlyTyrLeuIle 174
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383 GlyAspTPAsnThrThrLeuAlaCysPheValAlaIleLeuIleGlyLe 399
|||||
1453 GGAGACTGGAACACACATAGCCTGTTCGTAGCCATATTAATGGTTT 1502
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399 uCysLeuThrLeuLeuLeuAlaValPheLysLysAlaLeuProAlaL 416
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1503 GTGCCTTACATATTACTCCTTGCCATTTCAGAAAGCATTGCCAGCTC 1552
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416 euProIleSerIleThrPheGlyLeuIlePheTyrPheSerThrAspAsn 432
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seq_documentation_block:
; Sequence 1, Application US/08592541
; Patent No. 5986054
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/592,541
; FILING DATE:
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2791 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-592-541-1

alignment_scores:
Quality: 1461.00 Length: 466
Ratio: 3.938 Gaps: 8
Percent Similarity: 79.614 Percent Identity: 65.451

alignment_block:
US-08-509-3598-137 x US-08-592-541-1

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ATTORNEY/AGENT INFORMATION:
NAME: Han, William T
REGISTRATION NUMBER: 34,344
REFERENCE/DOCKET NUMBER: P50358
TELECOMMUNICATION INFORMATION:
TELEPHONE: 610-270-5219
TELEFAX: 610-270-5090
TELEX:
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1762 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-08-670-964-1

alignment_scores:
Quality: 1454.00 Length: 466
Ratio: 3.919 Gaps: 9
Percent Similarity: 79.614 Percent Identity: 65.451

alignment_block:
US-08-509-359B-137 x US-08-670-964-1 ..
Align seg 1/1 to: US-08-670-964-1 from: 1 to: 1762

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55 spGlyGluGluAspProArgTyrValCysSerGlyValPro..... 69
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277 GGCAGGAGCACAGCAGAGGAGCGCTTGGCCACCTGAGCCATTATCT 326
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70 ...GlyArgProGly..... 74
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75 LeuGluGluGluLeuThrLeuLysTyrGlyAlaLysHisValIleMetL 91
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158 ValLeuTyrLysTyrArgCysTyrLysPheIleHisGlyTrpIleLeuMe 174
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174 tSerSerLeuMetLeuLeuPheLeuThrTyrIleTyrLeuGlyGlu 191
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777 ATCTGGAATTTTGTGTGGTGAATGATTTCCCATTCACCTGGAAGGTCC 826
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224 oLeuValLeuGlnGlnAlaTyrLeuIleMetIleSerAlaLeuMetAla 241
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241 euValPheIleLysTyrLeuProGluTrpSerAlaTrpValIleLeuGly 257
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335 erTyrProGluValPheGluProProLeuThrGlyTyrProGly..... 349
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seq_documentation_block:

; Sequence 3, Application US/08670964

; Patent No. 6010874

; GENERAL INFORMATION:

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Date: Mar 18, 2000 11:52 PM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-1998 Compugen Ltd.

Command line parameters:

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-DB=EST -qfmt=fastap -SUFFIX=rst -GAPOP=12.000 -GAPEXT=4.000
-MINMATCH=0.100 -LOOPCL=0.000 -LOOPEXT=0.000 -GAPOP=4.500
-GAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -FGAPOP=6.000
-FGAPEXT=7.000 -YGAPOP=10.000 -YGAPEXT=0.500 -DELOP=6.000
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-LIST=45 -DOALIGN=200 -THR_SCORE=pct -ALIGN=15 -MODE=LOCAL
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Search information block:

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Query length: 448

Database: EST:*

Database sequences: 4538634

Database length: 1887831982

Search time (sec): 350.460000

score_list:

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gb_est31:AI675803	+ 1120.00	2273.18	7.5e-118	661	AI675803 w597a12.x1 NCI_CGAP.Pa
gb_est33:AI831581	+ 1118.00	2268.72	1.3e-117	680	AI831581 w39d04.x1 NCI_CGAP.Lu
gb_est21:AA993681	+ 1000.00	2028.34	3.3e-104	587	AA993681 ot97b02.s1 Soares_tot
gb_est39:AW131752	+ 916.00	1855.74	1.1e-91	598	AW131752 xf34b09.x1 NCI_CGAP.Br
gb_est17:AA602396	+ 890.00	1803.65	1.1e-94	538	AA602396 nc30d05.s1 NCI_CGAP.Br
gb_est35:AI829395	+ 877.00	1777.55	3.0e-90	513	AI829395 w64f10.x1 NCI_CGAP.Pa
gb_est33:AI765870	+ 870.00	1759.27	3.2e-89	707	AI765870 w66e09.x1 NCI_CGAP.Ki
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gb_est38:AAU56325	+ 835.50	1690.65	2.1e-85	591	AAU56325 zf53a03.s1 Soares_reti
gb_est25:AI276606	+ 824.00	1669.57	3.1e-84	481	AI276606 g171b07.x1 Soares.NhM
gb_est24:AI046485	+ 739.00	1494.79	1.7e-74	494	AI046485 uq70d05.y1 Sugano_mous
gb_est11:TO3796	+ 732.50	1481.90	8.9e-74	476	TO3796 IB913 Infant brain, Benc
gb_est23:AI097783	+ 704.00	1423.23	1.6e-70	483	AI097783 ue35g10.y1 Sugano_mous
gb_est36:AI927349	+ 704.00	1425.15	1.3e-70	413	AI927349 uq49f05.x1 NCI_CGAP.Lu
gb_est24:AI225996	+ 663.50	1354.72	1.1e-66	515	AI225996 u308a11.y1 Sugano_mous
gb_est11:AA260597	+ 663.50	1339.42	7.7e-66	511	AA260597 mx76d09.r1 Soares_mous
gb_est22:AI039365	+ 651.00	1316.98	1.4e-64	393	AI039365 ox40a06.s1 Soares_tot
gb_est44:AW177499	+ 650.00	1314.92	1.8e-64	465	AW177499 PM4-CT0155-270899-001
gb_est20:AA862334	+ 621.00	1255.76	2.3e-64	467	AA862334 oq94a07.s1 NCI_CGAP.Ki
gb_est19:AA782235	+ 612.00	1238.29	3.1e-60	352	AA782235 ai31c05.s1 Soares_para
gb_est14:AA451661	+ 591.00	1193.35	1.1e-57	409	AA451661 z343f07.r1 Soares_tot
gb_est15:AA537185	+ 555.00	1114.97	2.4e-53	591	AA537185 v445e03.r1 Soares_mous
gb_est11:AA268820	+ 543.50	1095.51	3.0e-52	421	AA268820 v501c10.r1 Soares_mous
gb_est25:AI324363	+ 539.50	1086.90	8.9e-52	435	AI324363 mr15d12.y1 Soares_mous
gb_est11:AA237206	+ 534.00	1074.67	4.3e-51	470	AA237206 mx18e01.r1 Soares_mous
gb_est10:AA162310	+ 495.00	991.87	1.8e-46	589	AA162310 mm44a07.r1 Beddington
gb_est5:N24576	+ 478.50	961.96	8.1e-45	426	N24576 yx72a04.s1 Soares_melan
gb_est11:AA265273	+ 475.00	956.11	1.7e-44	382	AA265273 mx91c12.r1 Soares_mous
gb_est5:H06456	+ 474.00	956.11	1.7e-44	323	H06456 y01b06.s1 Soares retina
gb_est26:AA817957	+ 474.00	950.50	3.5e-44	511	AA817957 UI-R-A0-ag-f-06-0-UI.s
gb_est10:AA144382	+ 473.50	950.53	3.5e-44	469	AA144382 mr15d12.r1 Soares_mous
gb_est25:AI287167	+ 472.00	950.48	3.5e-44	366	AI287167 u173h11.y1 Sugano_mous
gb_est2:RI14600	+ 470.50	949.59	4.0e-44	306	RI14600 yf34b10.r1 Soares_fetal
gb_est28:AI528670	+ 469.50	943.32	8.9e-44	432	AI528670 mr15d12.x1 Soares_mous
gb_est5:N27820	+ 456.00	913.88	3.9e-42	498	N27820 yx54h10.r1 Soares_melan
gb_est2:R05907	+ 455.50	918.53	2.1e-42	313	R05907 ye93h02.r1 Soares_fetal
gb_est11:AA231081	+ 423.00	843.04	3.4e-38	642	AA231081 mw11d11.r1 Soares_mous
gb_est4:HI19012	+ 421.50	842.57	3.6e-38	519	HI19012 ym44b02.r1 Soares_infant
gb_est11:AA210480	+ 417.50	834.50	1.0e-37	513	AA210480 mo86b03.r1 Beddington
gb_est18:AA673862	+ 413.00	824.06	3.9e-37	566	AA673862 v082h01.r1 Harstead md

gb_est4:H33787 + 412.00 831.38 1.5e-37 263 ! H33787 EST110123 Rat PC-12 c
gb_est11:AA200706 + 409.00 818.60 7.9e-37 452 ! AA200706 mu03f12.r1 Soares m
gb_est36:AI875963 + 409.00 823.02 4.5e-37 315 ! AI875963 uj53c08.y1 Sugano m

seq_name: gb_est36:AI925372

seq_documentation_block:
LOCUS AI925372 703 bp mRNA EST 02-SEP-1999
DEFINITION wn53d06.x1 NCI_CGAP_Lu19 Homo sapiens cDNA clone IMAGE:2449163 3'
similar to SW:PSN2_HUMAN P49810 PRESENILIN 2 ; , mRNA sequence.

ACCESSION AI925372

VERSION AI925372.1 GI:5661336

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 703)

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

On May 18, 1998 this sequence version replaced gi:3137011.

Contact: Robert Strausberg, Ph.D.

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Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

DNA Sequencing by: Greg Lennon, Ph.D.

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.lnl.gov/bbrp/image/image.html

Seq primer: -40UP from Gibco

High quality sequence from: 459.

Location/Qualifiers

FEATURES

source

1. 703

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:2449163"

/clone_lib="NCI_CGAP_Lu19"

/tissue_type="squamous cell carcinoma, poorly

differentiated (4 pooled tumors, including primary and

metastatic)"

/dev_stage="adult"

/lab_host="DH10B (phage-resistant)"

/note="Organ: lung; Vector: pT73D-Pac (Pharmacia) with a

modified polylinker; 1st strand cDNA was prepared from

pooled lung tumor tissue, and was then primed with a Not I

- oligo(dT) primer. Double-stranded cDNA was ligated to

Eco RI adaptors (Pharmacia), digested with Not I and

cloned into the Not I and Eco RI sites of the modified

pT73 vector. Library went through one round of

normalization. Library constructed by Bento Soares and M.

Fatima Bonaldo."

BASE COUNT 138 a 203 c 177 g 184 t 1 others

ORIGIN

alignment_scores:

Quality: 1155.00

Ratio: 5.088

Percent Similarity: 97.425

Percent Identity: 97.425

alignment_block:

US-08-509-359B-137 x AI925372

Align seg 1/1 to: AI925372 from: 1 to: 703

79 LeuThrLeuLysTyrGlyAlaLysHsValleMetLeupheValProva 95
|||||

195 rAsnValAlaMetAspTyrProThrLeuLeuLeuThrValTtpAsnPhg 212

352 CAATGGCCATGGACTACCCACCCTTGTGCTGACTGTCTGGAACTTCG 401

212 lyAlaValGlyMetValCysIleHisTrpLysGlyProLeuValLeuGln 228

402 GGGCAGTGGGCATGTGTGATCCACTGGAAGGCCCTCTGGTGTGCAG 451

229 GlnAlaTyrLeuIleMetIleSerAlaLeuMetAlaLeuValPheIleLy 245

452 CAGGCTACCTCATCATGATCATGTCGCTGATGGCCCTAGTGTTCATCAA 501

245 sTyrLeuProGluTtpSerAlaTtpValIleLeuGlyAlaIleSerValT 262

502 GTACCTCCAGAGTGGTCCGGTGGGTGATCTCGGCCCATCTCTGTGT 551

262 yAspLeuValAlaValLeuCysProLysGlyProLeuArgMetLeuVal 278

552 ATGATCTCGTGGCTGTGTGTGCCANAGGCCCTCTGAGAAATGCTGTA 601

279 GluThrAlaGlnGluArgAsnGluProIlePheProAlaLeuIleTyrSe 295

602 GAAACTGCCAGGAGAAATGAGCCCATATTCCTGCTTCATATACTC 651

295 rSerAlaMet 298

652 ATCTGNCATG 661

seq_name: gb_est35:AI831581

seq_documentation_block:

LOCUS AI831581 680 bp mRNA EST 26-AUG-1999
DEFINITION wJ39Q04.x1 NCI-CGAP_Lu19 Homo sapiens cDNA clone IMAGE:2405191 3',
similar to SW:PSN2_HUMAN P49810 PRESENILIN 2 ;, mRNA sequence.

ACCESSION AI831581

VERSION AI831581.1 GI:5452252

KEYWORDS EST.

SOURCE human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 680)

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

On Dec 20, 1995 this sequence version replaced gi:1130797.

Contact: Robert Strausberg, Ph.D.

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Email: Robert.Strausberg@nih.gov

Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

DNA Sequencing by: Greg Lennon, Ph.D.

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Seq primer: -40UP from Gibco

High quality sequence stop: 456.

FEATURES

source

1. .680

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:2405191"

/clone_lib="NCI-CGAP_Lu19"

/tissue_type="squamous cell carcinoma, poorly

differentiated (4 pooled tumors, including primary and

metastatic)"

/dev_stage="adult"

/lab_host="DH10B (phage-resistant)"

/note="Organ: lung; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from pooled lung tumor tissue, and was then primed with a Not I - oligo(drf) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 137 a 191 c 173 g 178 t 1 others
ORIGIN

alignment_scores:

Quality: 1118.00 Length: 226

Ratio: 5.059 Gaps: 0

Percent Similarity: 97.788 Percent Identity: 96.903

alignment_block:

US-08-509-359B-137 x AI831581 ..

Align seg 1/1 to: AI831581 from: 1 to: 680

79 LeuThrLeuLysTyrGlyAlaLysHisValIleMetLeuPheValProVa 95

3 CTGACCCCTCAAAATACATGATCGAAGCACGATGATCTGTGTGCGCTGT 52

95 lThrLeuCysMetIleValValAlaThrIleLysSerValArgPheT 112

53 CACTCTGTGCATGATCGTGTGTAGCCACCATCAAGTCTGTGCGCTTCT 102

112 YThrGluLysAsnGlyGlnLeuIleTyrThrProPheThrGluAspThr 128

103 ACACAGAGAAGATGACAGCTCATCTACAGCCATCTACTGAGGACACA 152

129 ProSerValGlyGlnArgLeuLeuAsnSerValLeuAsnThrLeuIle 145

153 CCCTCGTGGGCCAGCGCTCTCAACTCCGTGCTGAACACCCCTCATCAT 202

145 tIleSerValIleValValMetThrIlePheLeuValValLeuTyrLysT 162

203 GATCAGCGTCATCGTGGTATTACCATCTCTTGTGTGCTCTACAAGT 252

162 YArgCysTyrLysPheIleHisGlyTrpLeuIleMetSerSerLeuMet 178

253 ACCGCTGCTACAAAGTTCATCCATGGCTGGTGTGATCATGCTTCACTGATG 302

179 LeuLeuPheLeuPheThrIleTyrLeuGlyGluValLeuLysThrTy 195

303 CTGCTGTTCCTCTTCACTATATCTACCTTGGGGAAGTCTCAAGACCTA 352

195 rAsnValAlaMetAspTyrProThrLeuLeuLeuThrValTtpAsnPhg 212

353 CAATGGGCATGGACTACCCACCCCTCTGTGACTGTCTGGAACTTCG 402

212 lyAlaValGlyMetValCysIleHisTrpLysGlyProLeuValLeuGln 228

403 GGGCAGTGGGCATGGTGTGCATCCACTGGAAGGCCCTCTGGTGTGCAG 452

229 GlnAlaTyrLeuIleMetIleSerAlaLeuMetAlaLeuValPheIleLy 245

453 CAGGCTTACCTCATCATGATGATGCGCTCATGGCCCTAGTGTTCATCAA 502

245 sTyrLeuProGluTtpSerAlaTtpValIleLeuGlyAlaIleSerValT 262

503 GTACCTCCAGAGTGGTCCGGTGGGTGATCTACGCGCCATCTCTGTGT 552

262 YrAspLeuValAlaValLeuCysProLysGlyProLeuArgMetLeuVal 278

553 ATGATCTCGTGGCTGTGTGTACCATAGGCCCTCTGAGAAATGCTGGTA 602

279 GluThrAlaGlnGluArgAsnGluProIlePheProAlaLeuIleTyrSe 295

High quality sequence stop: 397.

FEATURES

Location/Qualifiers
1..597
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2619929"
/tissue_type="NCI_CGAP_Brn50"
/tissue_type="medulloblastoma"
/lab_host="DH10B (phage resistant)"
/note="Organ: Brain; Vector: pT73D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from medulloblastoma tumor tissue, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. This library is normalized. Library constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 119 a 170 c 147 g 161 t

alignment_scores:

Quality: 916.00 Length: 199
Ratio: 4.796 Gaps: 0
Percent Similarity: 95.980 Percent Identity: 92.462

alignment_block:

US-08-509-359B-137 x AW131752

Align seg 1/1 to: AW131752 from: 1 to: 597

79 LeuThrLeuLysTyrGlyAlaLysHisValIleMetLeuPheValProva 95
|||||
2 CTGACCCCTCAATACTG.TCGAAGCACGTGATCATGCTGTTCGCCCTGT 50
95 lThrLeuCysMetIleValValAlaThrIleLysSerValArgPheT 112
51 CACCTGTGTCATGATCGTGGTGGTAGCCACCATCAAGTCTGCGCCTCT 100
112 yrThrGlyAsnGlyGlnLeuIleTyrThrProPheThrGluAspThr 128
101 ACACAGAGAAGAAATGGACAGCTCATCTACAGCCATTCTCAGTGAGACACA 150
129 ProSerValGlyGlnArgLeuLeuAsnSerValLeuAsnThrLeuIleMe 145
151 CCCTCGGTGGGCCAGCGCTCTCTCAACTCCGTGCTGAACCCCTCATCAT 200
145 tIleSerValIleValValMetThrIlePheLeuValValLeuTyrLysT 162
201 GATCAGCTCATCGTGGTATGACCATCTTCTGTGGTGTCTCTCAAGT 250
162 yrArgCysTyrLysPheIleHisGlyTrpLeuIleMetSerSerLeuMet 178
251 ACCGTGCTACAAGTTTCATCCATGCTGCTGATCATGCTCTCATCTGATG 300
179 LeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluValLeuLysThrTy 195
301 CTGCTGTTCCTCTTCCACCTATATCTACCTTGGGAAGTGCCTCAAGACCTA 350
195 rAsnValAlaMetAspTyrProThrLeuLeuLeuThrValTrpAsnPhG 212
351 CAATGTGCCCATGGACTACCCACCCCTTGTGACTGTCTGGAACCTTCG 400
212 lyAlaValGlyMetValCysIleHisTrpLysGlyProLeuValLeuGln 228
401 GGGCAGTGGGATGGTGTGCATCCACTCGAAGGGCCCTCTGGTGTGCGAG 450
229 GlnAlaTyrLeuIleMetIleSerAlaLeuMetAlaLeuValPheIleTy 245
451 CACGCCTACCTCATCATCATCAGTCGCTCATGTCCTATTGTTTCATCAA 500
245 sTyrLeuProGlnTrpSerAlaTrpValIleLeuGlyAlaIleSerValT 262
| |||||

501 GATCCTACCAAGTGTACGCGTGGTGCATCTGACGCCATATCTGTGT 550

262 yrAspLeuValAlaValLeuCysProLysGlyProLeuArgMetLeu 277

551 ATGATCTCGTCCCTTGGCTGTGTACCAATGGCCTGTGTGAATGCTG 597

seq_name: gb_est17:AA602396

seq_documentation_block:

LOCUS AA602396 538 bp mRNA EST 08-OCT-1997
DEFINITION nc30805.s1 NCI_CGAP_Pr22 Homo sapiens cDNA clone IMAGE:1102185 3'
Similar to TR:G1244640 G1244640 PRESENILIN I-374. ; mRNA sequence.
ACCESSION AA602396
VERSION AA602396.1 GI:2436374
KEYWORDS EST.
SOURCE human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT On Sep 19, 1997 this sequence version replaced gi:1517341.

Contact: Robert Strausberg, Ph.D.

Tel: (301) 496-1550

Email: Robert.Strausberg@nih.gov

Tissue Procurement: Michael J. Brownstein, M.D., Ph.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 1338 Std Error: 0.00

Seq primer: -40ml3 fwd. ET from Amersham

High quality sequence stop: 404.

FEATURES

source

1..538
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1102185"
/clone_lib="NCI_CGAP_Pr22"
/sex="male"
/tissue_type="normal prostate"
/lab_host="DH10B"
/note="Organ: prostate; Vector: pT73D-Pac (Pharmacia)
with a modified polylinker; 1st strand cDNA was prepared
from normal prostate bulk tissue, and was then primed with
a Not I - oligo(dT) primer. Double-stranded cDNA was
ligated to Eco RI adaptors (Pharmacia), digested with Not
I and cloned into the Not I and Eco RI sites of the
modified pT73 vector. Library is normalized, and was
constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 101 a 155 c 138 g 143 t 1 others

ORIGIN

alignment_scores:

Quality: 890.00 Length: 179
Ratio: 5.028 Gaps: 0
Percent Similarity: 98.883 Percent Identity: 98.324

alignment_block:

US-08-509-359B-137 x AA602396

Align seg 1/1 to: AA602396 from: 1 to: 538

94 ProValThrLeuCysMetIleValValAlaThrIleLysSerValAr 110

|||||

DEFINITION wh66e09.x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:2385736 3' similar to SW:PSN2_HUMAN P49810 PRESENILIN 2 ;, mRNA sequence.

ACCESSION A1765870

VERSION A1765870.1 GI:5232379

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

REFERENCE 1 (bases 1 to 707)

AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

JOURNAL Unpublished (1997)

COMMENT On Feb 17, 1998 this sequence version replaced gi:2889754.

Contact: Robert Strausberg, Ph.D.

Tel: (301) 496-1550

Email: Robert.Strausberg@nih.gov

Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone Distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html

Seq primer: -40UP from Gibco

High quality sequence stop: 466.

Location/Qualifiers

1..707

FEATURES

source

1..707

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:2385736"

/clone.lib="NCI_CGAP_Kid11"

/lab.host="DH10B"

/note="Organ: kidney; Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; Plasmid DNA from the normalized library NCI_CGAP_Kid3 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneids 1323376-1323911, 1456007-1456775, and 1500552-1502855). Subtraction by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 177 a 197 c 182 g 150 t 1 others

ORIGIN

alignment_scores

Quality: 870.00 Length: 186

Ratio: 4.807 Gaps: 0

Percent Similarity: 97.312 Percent Identity: 95.161

alignment_block

US-08-509-359b-137 x A1765870/rev ..

Align seg 1/1 to reverse of: A1765870 from: 1 to: 707

263 AspLeuValAlaValLeuCysProLysGlyProLeuArgMetLeuValG1 279

||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||

707 GATCTGGGGCGTGTGTGTCTCCAAAGGCGCTTCTGAGATGCTGTAGA 658

279 uThrAlaGlnGluArgAsnGluProIlePheProAlaLeuIleTyrSers 296

||||| :||||| :||||| :||||| :||||| :||||| :||||| :|||||

657 AACTGCCAGAGAGAAATGAGCCCATATCTCCCTGCCCTGATATA. NCAT 609

296 exAlaMetValTrpThrValGlyMetAlaLysLeuAspProSerSergln 312

||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||

608 CTGCCATGTTGGACGGTGGCATGGGAAGCTGGACCCCTCC.TCTCAG 560

313 GlyAlaLeuGlnLeuProTyrAspProGluMetGluGluAspSerTyrAs 329

||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||

559 GGTGCCCTCCAGTCCCTACGACCCGGAGATGGAGAGACTCCTATGA 510

329 pSerPheGlyGluProSerTyrProGluValPheGluProProLeuThrG 346

||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||

509 CAGTTTTGGGGAGCCTTCATACCCCGAAGTCTTTGAGCCTCCCTTGACTG 460

346 lYrProGlyGluGluLeuGluGluGluGluGluGluGluGluGluGlu 362

||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||

459 GCTACCCAGGGAGGAGCTGGAGGAAGAGGAGGAGGCGCTGAAGCTT 410

363 GlyLeuGlyAspPheIlePheTyrSerValLeuValGlyLysAlaAla 379

||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||

409 GGCTCTGGGAGCTTCATCTTCTACAGTGTCTGTGGCAGGCGGCTGC 360

379 aThrGlySerGlyAspTrpAsnThrThrLeuAlaCysPheValAlaLe 396

||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||

359 CACGGGCGAGCGGAGCTGGAATACACGCTGGCCTGCTTCGTGGCCATCC 310

396 euIleGlyLeuCysLeuThrLeuLeuLeuLeuAlaValPheLysLysAla 412

||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||

309 TCATTGGCTTGTGTCTGACCTCTCTGCTGTCTGTGTGTTCAGAAGGCG 260

413 LeuProAlaLeuProIleSerIleThrPheGlyLeuIlePheTyrPheSe 429

||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||

259 TGGCCCCCCTCCCATTCATCCAGTTCGGGCTCATCTTTTACTTCTC 210

429 rThrAspAsnLeuValArgProPheMetAspThrLeuAlaSerHisGlnL 446

||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||

209 CACGGACAACCTGGTGGCGCGGTTCATGGACACCTGGCCTCCCATCAGC 160

446 euTyrIle 448

|||||

159 TCTACATT 152

seq_name: gb_est44:AW212769

seq_documentation_block:

LOCUS AW212769 555 bp mRNA EST 03-DEC-1999

DEFINITION u066e03.x1 NCI_CGAP_Mam1 Mus musculus cDNA clone IMAGE:2647516 3' similar to gb:L42177 Mus musculus S182 protein mRNA, complete cds (MOUSE);, mRNA sequence.

ACCESSION AW212769

VERSION AW212769.1 GI:6518888

KEYWORDS EST.

SOURCE house mouse.

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 555)

AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

JOURNAL Unpublished (1997)

COMMENT

On Jul 7, 1999 this sequence version replaced gi:5406916.

Other_ESTs: u066e03.y1

Contact: Robert Strausberg, Ph.D.

Tel: (301) 496-1550

Email: Robert.Strausberg@nih.gov

Tissue Procurement: Gilbert Smith, Ph.D.

cDNA Library Preparation: Life Technologies, Inc.

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Washington University Genome Sequencing Center

Clone Distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

MG1:1027968

Seq primer: -40UP from Gibco

High quality sequence stop: 421.

Location/Qualifiers

1..555

/organism="Mus musculus"

FEATURES

source


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3 CCTGTCACTCTGTGCATGTCGGTGTAGCCACCATCAAGTCTGGCG 52
110 gPheTyrThrGluLysAsnGlyGlnLeuLeuTyrThrProPheThrGluA 127
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53 CTTCTACACAGAGAAGATGGACAGCTCATCTACACGCCATTCACCTGAGG 102
127 sPThrProSerValGlyGlnArgLeuLeuAsnSerValLeuAsnThrLeu 143
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103 ACACACCTCGTGGGCCAGCGCTCCTCACTCCGTGTGAACACCTC 152
144 IleMetIleSerValIleValValMetThrIlePheLeuValValLeuTy 160
153 ATCATGATCAGCGTCATCGTGGTATGACCATCTTCTGGTGTGCTCTA 202
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210 nPheGlyAlaValGlyMetValCysIleHisTrpLysGlyProLeuVal 227
353 CTTGGGGGAGTGGGATGGTGTGATCCACTGGAAGGGCCCTCTGGTGC 402
227 euGlnGlnAlaTyrLeuIleMetIleSerAlaLeuMetAlaLeuValPhe 243
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244 IleLysTyrLeuProGluTrpSerAlaTrpValIleLeuGlyAlaIleSe 260
453 ATCAAGTACTC.CCAGAGTGGTCCGGTGGTGCATC...TGGCGGCCANC 498
260 rValTyrAspLeuValAlaValLeuCysProLysGlyProLeuArgMetL 277
499 TCTGTGTATGATCCGTNGTGTGTGTGTCCAAAGGCT...CTGAGAATGT 545
277 euValGluThrAlaGlnGluArgAsnGluPro 287
546 GG...TAGAANTGCCAGGAGAGAATGAGCCA 574

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seq_name: gb_est25:A1276606

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seq_documentation_block: 481 bp mRNA EST 29-JAN-1999
LOCUS A1276606
DEFINITION gi71h07.x1 Soares_NHMPu_s1 Homo sapiens cDNA clone IMAGE:1877821
3', similar to SW:PSN2_HUMAN P49810 PRESENILIN 2 ;, mRNA sequence.
ACCESSION A1276606
VERSION A1276606.1 GI:3898880
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 481)
NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
On Jan 19, 1998 this sequence version replaced gi:2285413.
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
Insert Length: 1769 Std Error: 0.00
Seq primer: -400P from Gibco

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High quality sequence stop: 448.

Location/Qualifiers

1..481

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:1877821"

/clone_lib="Soares_NHMPu_s1"

/tissue_type="Pooled human melanocyte, fetal heart, and pregnant uterus"

/lab_host="DH10B"

/note="Organ: mixed (see below); Vector: pT7R3D-Pac (Pharmacia) with a modified polylinker; Site.1: Not I; Site.2: Eco RI; Equal amounts of plasmid DNA from three normalized libraries (melanocyte 2NBHM, pregnant uterus NDHPU, and fetal heart NDHH19W) were mixed, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from pools of 5,000 clones made from the same 3 libraries. The pools consisted of I.M.A.G.E. clones 260232-265223, 340488-345479, and 484488-489479."

BASE COUNT 100 a 138 c 119 g 124 t

ORIGIN

alignment_scores:

Quality: 824.00 Length: 160

Ratio: 5.150 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-08-509-359B-137 x A1276606 ..

Align seg 1/1 to: A1276606 from: 1 to: 481

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77  GUGLULeuThrLeuLysTyrGlyAlaLysHisValIleMetLeuPheVa 93
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2  GAGAGCTGACCTCAANTAGGAGGAGCATGTGATCATGCTGTTGT 51
93  lProValThrLeuCysMetIleValValAlaThrIleLysSerValA 110
|||||
52  GCCTGTCACTCTGTGCATGATCGTGGTGTAGCCACCATCAAGTCTG 101
110  rPheTyrThrGluLysAsnGlyGlnLeuLeuTyrThrProPheThrGlu 126
|||||
102  GCTTCTACACAGAGAGATGGACAGCTCATCTACAGCCATTCAC 151
127  AspThrProSerValGlyGlnArgLeuLeuAsnSerValLeuAsnThrLe 143
|||||
152  GACACACCTCGGTGGGCCAGCGCTCTCAACTCCGTGTGAACACCT 201
143  uIleMetIleSerValIleValValMetThrIlePheLeuValValLeu 160
202  CATCATGATCAGCGTCATCGTGGTATGACCATCTTCTTGGTGTGCTCT 251
160  yrlLysTyrArgCysTyrLysPheIleHisGlyTrpLeuIleMetSerSer 176
|||||
252  ACAAGTACCCTGTGTACAAAGTTCATCCATCGCTGTTGATCATGCTTCA 301
177  LeuMetLeuLeuPheLeuPheThrTyrIleTyrLeuGlyGluValLeuLy 193
|||||
302  CTGATGCTGTGTTCTCTTACCATATATCTACCTTGGGAAGTGTCTCAA 351
193  sThrTyrAsnValAlaMetAspTyrProThrLeuLeuLeuThrValTPA 210
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352  GACCTACAATGTGGCCATGAGTACCCACCTCTTGTGACTGTCTGGA 401
210  snPheGlyAlaValGlyMetValCysIleHisTrpLysGlyProLeuVal 226
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seq_documentation_block:			
LOCUS	AI097783	483 bp	EST
DEFINITION	ue35g10.y1 Sugano mouse liver mliia Mus musculus cDNA clone		
IMAGE:	1482402 5' similar to gb:L42177 Mus musculus S182 protein mRNA, complete cds (MOUSE);, mRNA sequence.		

GenCore version 4.5
Copyright (c) 1993 - 1998 Compugen Ltd.

OM protein - protein search, using sw model

Run on: March 20, 2000, 04:21:11 ; Search time 35.25 Seconds
(without alignments)
301.032 Million cell updates/sec

Title: US-08-509-359B-137
Perfect score: 2336
Sequence: 1 MLTFMASDSEEEVCDERTSL.....STDNLVRFMDTLASHQLYI 448

Scoring table: BLOSUM62

Searched: 188963 seqs, 23686106 residues

Database : A_Geneseq_36:*

Word size : 0

Number of hits that pass the threshold : 188963

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	2336	100.0	448	1	W05762	Human presenilin-2
2	2336	100.0	448	1	W11321	Human AD4 protein.
3	2336	100.0	448	1	W23967	Human presenilin-2
4	2332	99.8	448	1	W05763	Presenilin-2 M339V
5	2331	99.8	448	1	W05765	Presenilin-2 I420T
6	2327	99.6	448	1	W05764	Presenilin-2 N141I
7	2320.5	99.3	447	1	W28508	Full AD4/AD3LP seq
8	2136	91.4	414	1	W05766	Presenilin-2 delta
9	1623.5	69.5	376	1	W28506	AD4/AD3LP sequence
10	1468	62.8	467	1	W05735	Murine presenilin-1
11	1468	62.8	467	1	W23966	Mouse presenilin-1
12	1467	62.8	467	1	W05733	Presenilin-1-1. Ne
13	1467	62.8	467	1	W41430	PS1/467 protein. D
14	1467	62.8	467	1	W05755	Human presenilin-1
15	1464	62.7	467	1	W05754	Presenilin-1-1 L28
16	1464	62.7	467	1	W05758	Presenilin-1-1 L39
17	1464	62.7	467	1	W05737	Presenilin-1-1 V82
18	1464	62.7	467	1	W05746	Presenilin-1-1 I21
19	1463	62.6	467	1	W05754	Presenilin-1-1 A28
20	1463	62.6	467	1	W05736	Presenilin-1-1 A79
21	1463	62.6	467	1	W05747	Presenilin-1-1 I23
22	1463	62.6	467	1	W05749	Presenilin-1-1 A26
23	1463	62.6	467	1	W41431	Mouse PSI/467 prot
24	1462	62.6	467	1	W05738	Presenilin-1-1 V96
25	1462	62.6	467	1	W05739	Presenilin-1-1 Y11
26	1462	62.6	467	1	W05741	Presenilin-1-1 I14
27	1462	62.6	467	1	W05748	Presenilin-1-1 A24
28	1462	62.6	467	1	W27176	Human S182 gene, P
29	1461	62.5	467	1	W05753	Presenilin-1-1 E28
30	1461	62.5	467	1	W05757	Presenilin-1-1 G38
31	1461	62.5	467	1	W05740	Presenilin-1-1 M13
32	1461	62.5	467	1	W05742	Presenilin-1-1 M14
33	1460	62.5	467	1	W05744	Presenilin-1-1 L17
34	1460	62.5	467	1	W56770	Homo sapiens PS-1.
35	1459	62.5	467	1	W05752	Presenilin-1-1 P26
36	1458	62.4	467	1	W05743	Presenilin-1-1 H16
37	1458	62.4	467	1	W05745	Presenilin-1-1 G20
38	1457	62.4	467	1	W05751	Presenilin-1-1 P26
39	1456	62.3	467	1	W05759	Presenilin-1-1 C41

40 1455 62.3 467 1 W05750 Presenilin-1-1 C26
41 1454 62.2 467 1 W11839 Human early onset
42 1454 62.2 463 1 W11840 Early onset Alzheimer
43 1448 62.0 463 1 W05734 Presenilin-1-2. Ne
44 1448 62.0 463 1 W23965 Human presenilin-1
45 1446 61.9 463 1 W42375 Human presenilin I

ALIGNMENTS

RESULT 1

W05762

ID W05762 standard; Protein; 448 AA.

AC W05762;

DT 25-JUL-1997 (first entry)

DE Human presenilin-2.

KW Presenilin-2; human; hpsl-1; hpsl-2; PS-2; integral membrane protein; AD;

KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;

KW depression; antibody; gene expression modulator; therapy.

OS Homo sapiens.

PN W09634099-A2.

PD 31-OCT-1996.

PF 29-APR-1996; CA0263.

PR 28-APR-1995; US-431048.

PR 28-JUN-1995; US-496841.

PR 31-JUL-1995; US-509359.

PA (HSCR-) HSC RES & DEV LP.

PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.

PI Fraser PE, Rommens JM, St George-Hyslop PH;

DR WPI: 96-497631/49.

DR N-FSDB; T40031.

PT New presenilin genes - useful for diagnosis, therapy and drug

PT screening of familial Alzheimer's disease, cerebral disorders, etc.

PS Claim 4; Page 148-150; 178pp; English.

CC This sequence represents the wild type human presenilin-2.

CC W05733 and W05734 represent the two different forms of wild type human

CC presenilin-1 (PS-1). The form represented by W05734 results from

CC alternate splicing of the genomic DNA sequence. W05735 represents the

CC coding sequence for wild type mouse PS-1. The presenilins are a family of

CC highly conserved integral membrane proteins with a common structural

CC motif, common alternate splicing patterns, and common mutational hot spot

CC regions. Mutations in PS genes are implicated in familial Alzheimer's

CC disease (AD) and possibly other diseases such as cerebral haemorrhage,

CC schizophrenia, depression etc., so detection of mutations in the DNA

CC encoding these sequences can be used for diagnosis of these diseases.

CC These proteins, or vectors that express them or containing antisense

CC sequences, antibodies selective for mutant forms of these proteins (such

CC as W05736) and modulators of PS gene expression are potentially useful

CC for treatment of AD etc. Transgenic animals are useful as models for drug

CC screening. The antibodies can also be used e.g. for affinity purification

CC and in immunoassays.

SQ Sequence 448 AA;

Query Match 100.0%; Score 2336; DB 1; Length 448;

Best Local Similarity 100.0%; Pred. No. 7.3e-237;

Matches 448; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 MLTFMASDSEEEVCDERTSLMSAESPTPRSCQGRGPDGENTAQRSGNEDEGEDP 60

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Db 61 DRYVCSGVPRPPGLEEEETLKYGAHVIMLFVPVTLCMVVVATIKSVRFYKNGQLI 120

QY 121 YPPTFEDTPSVQGRLLNSVINTLIMISIVVMTIFLVLYKYKCYKFIHGWLIMSSLM 180

Db 121 YPPTFEDTPSVQGRLLNSVINTLIMISIVVMTIFLVLYKYKCYKFIHGWLIMSSLM 180

QY 181 FLFTYIYGEVKTNYVNDYPTLLTWNFGVGVCHWKGPLVLOQAYLIMISALMA 240

Db 181 FLFTYIYGEVKTNYVNDYPTLLTWNFGVGVCHWKGPLVLOQAYLIMISALMA 240

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 FT /note= "hydrophilic loop"
 FT 201. 218
 FT /label= TM4
 FT /note= "transmembrane domain 4"
 FT 219. 224
 FT /label= TM4-5
 FT /note= "hydrophilic loop"
 FT 225. 244
 FT /label= TM5
 FT /note= "transmembrane domain 5"
 FT 245. 249
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 FT /note= "hydrophilic loop"
 FT 250. 268
 FT /label= TM6
 FT /note= "transmembrane domain 6"
 FT 269. 387
 FT /label= TM6-7
 FT /note= "hydrophilic loop"
 FT 388. 409
 FT /label= TM8
 FT /note= "transmembrane domain 8"
 FT 141
 FT /note= "Asn141lle mutation site (Claim 19)"
 FT 239
 FT /note= "Met239Val mutation site (Claim 19)"
 FT 420
 FT /note= "Ile420Thr mutation site"
 FT W09801549-A2.
 FT 15-JAN-1998.
 FT 04-JUL-1997. CA0475.
 FT 02-JAN-1997. US-034590.
 FT 05-JUL-1996. US-021673.
 FT 12-JUL-1996. US-021700.
 FT 08-NOV-1996. US-029895.
 FT (HSCR-) HSC RES & DEV LP.
 FT (UTOR) UNIV TORONTO GOVERNING COUNCIL.
 FT Fraser PE, Rommens JM, St George-Hyslop PH;
 FT WPI; 98-286355/25.
 FT N-PSDB; V04669.
 FT New isolated mutant presenilin-1 genes - useful for developing
 FT products for use in detection, diagnosis and therapy of Alzheimer's
 FT disease and for drug screening
 FT Claim 19; Page 203-204; 238pp; English.
 FT This polypeptide comprises human presenilin-2 (hPS2). Its amino
 FT acid sequence was deduced from an isolated cDNA clone (see V04669).
 FT Human and murine presenilin-1 sequences are also provided (see
 FT W23964-66). Mutations in the PS-1 and PS-2 genes are linked to
 FT the development in humans of forms of familial Alzheimer's disease
 FT (FAD) and may be causative of other disorders, e.g. cognitive,
 FT intellectual, neurological or physiological disorders such as
 FT cerebral haemorrhage, schizophrenia, depression, mental retardation
 FT and epilepsy. Use of the nucleic acids and proteins comprising or
 FT derived from the presenilins is made in screening and diagnosing
 FT FAD, identifying and developing therapeutics for treatment of FAD,
 FT and in producing cell lines and transgenic animals useful as models
 FT of FAD. Methods for identifying substances that bind to, or
 FT modulate the activity of a presenilin protein, and methods for
 FT identifying substances that affect the interaction of a
 FT presenilin-interacting protein with a presenilin protein are also
 FT disclosed. 448 AA;

Query Match 100.0%; Score 2336; DB 1; Length 448;
 Best Local Similarity 100.0%; Pred. No. 7.3e-237;
 Matches 448; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MLTFMADSEEVCDERTSLMSAESPTPRSCQEGRGQPEDGENTAQRWSQNEEDGEDP 60
 DB 1 MLTFMADSEEVCDERTSLMSAESPTPRSCQEGRGQPEDGENTAQRWSQNEEDGEDP 60

QY 61 DRYVCSGVPGRPPGLEELTLKYGAKHVIMLFVPTLCMIVVVAIKSVRFYTERNGQLI 120
 DB |||||||
 QY 121 YTPFTEDPSVQORLLNSVLNTLIMISIVVMTIFLVLYKYRCYKFHGWILMSMLL 180
 DB |||||||
 QY 181 FLFTYIYLGEVLKTYNVAMDYPTLLLTWNFGAVGMVCIHMKGPLVLOQAYLIMSALMA 240
 DB |||||||
 QY 241 LVFIKYLPEWSAWILGAISVYDLVAVLCPKGRLMVLTAQERNEPFPALIISSAMVW 300
 DB |||||||
 QY 301 TVGMAKLDPSSQALQLPYDPMEEDSDYDFGEPSPYEVFEPPLTGYGGEELSEEEERG 360
 DB |||||||
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 DB |||||||
 QY 421 TFGLIIFYESTDNLVRPFMDTLASHOLYI 448
 DB |||||||
 QY 421 TFGLIIFYESTDNLVRPFMDTLASHOLYI 448
 DB |||||||

RESULT 4
 W05763
 ID W05763 standard; Protein; 448 AA.
 AC W05763; 1997 (first entry)
 DT 25-JUL-1997 (first entry)
 DE Presenilin-2 M239V mutation.
 KW Presenilin-2; human; hPS1-1; hPS1-2; PS-2; integral membrane protein; AD;
 KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;
 KW depression; antibody; gene expression modulator; therapy; mutain.
 OS Homo sapiens.
 FH Key Location/Qualifiers
 FT modified_site 239 /label= M239V
 FT W05634099-A2.
 PD 31-OCT-1996.
 PF 29-APR-1996; CA0263.
 PR 28-APR-1995; US-431048.
 PR 28-JUN-1995; US-496841.
 PR 31-JUL-1995; US-509359.
 PA (HSCR-) HSC RES & DEV LP.
 PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
 PI Fraser PE, Rommens JM, St George-Hyslop PH;
 DR WPI; 96-497631/49.
 DR New presenilin genes - useful for diagnosis, therapy and drug
 DR screening of familial Alzheimer's disease, cerebral disorders, etc.
 PT Claim 4; Page -; 178pp; English.
 PS W05763-W05766 represent mutated versions of the human presenilin-2
 CC protein (see W05762 for wild type sequence). The presenilins are a family
 CC of highly conserved integral membrane proteins with a common structural
 CC motif, common alternate splicing patterns, and common mutational hot spot
 CC regions. Mutations in PS genes are implicated in familial Alzheimer's
 CC disease (AD) and possibly other diseases such as cerebral haemorrhage,
 CC schizophrenia, depression etc., so detection of mutations in the DNA
 CC encoding the wild type sequences can be used for diagnosis of these
 CC diseases. The wild type proteins, or vectors that express them or
 CC containing antisense sequences, antibodies selective for these mutant
 CC forms of the proteins and modulators of PS gene expression are
 CC potentially useful for treatment of AD etc. Transgenic animals are useful
 CC as models for drug screening. The antibodies can also be used e.g. for
 CC affinity purification and in immunoassays.
 SQ Sequence 448 AA;

Query Match 99.8%; Score 2332; DB 1; Length 448;

CC regions. Mutations in PS genes are implicated in familial Alzheimer's disease (AD) and possibly other diseases such as cerebral haemorrhage, schizophrenia, depression etc., so detection of mutations in the DNA encoding the wild type sequences can be used for diagnosis of these diseases. The wild type proteins, or vectors that express them or CC containing antisense sequences, antibodies selective for these mutant CC forms of the proteins and modulators of PS gene expression are potentially useful for treatment of AD etc. Transgenic animals are useful CC as models for drug screening. The antibodies can also be used e.g. for CC affinity purification and in immunoassays.

SQ Sequence 448 AA;

Query Match 99.6%; Score 2327; DB 1; Length 448;
Best Local Similarity 99.8%; Pred. No. 6.4e-236;
Matches 447; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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DB 1 MLTFMADSEEEVCDERTSLMSAESPTRSCQEGRGQPEDGENTAQRSENEDEEDP 60

QY 61 DRYVCSGVPGRPPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120
DB 61 DRYVCSGVPGRPPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120

QY 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSMLL 180
DB 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSMLL 180

QY 181 FLFTYIYLGVLKTYNVAMDYPTLLTWNFGAVGMVCIHWKGPLVLOQAYLIMISALMA 240
DB 181 FLFTYIYLGVLKTYNVAMDYPTLLTWNFGAVGMVCIHWKGPLVLOQAYLIMISALMA 240

QY 241 LVFTKYLPEWSAWILGAISSYDVLVAVLCPKGLRMLVETAQERNEPIFPALIISSAMVW 300
DB 241 LVFTKYLPEWSAWILGAISSYDVLVAVLCPKGLRMLVETAQERNEPIFPALIISSAMVW 300

QY 301 TVGMAKLDPSSQALQLPYDPEMEEDSDYSGEPSYPEVFPPLTGYGPELEEEERGV 360
DB 301 TVGMAKLDPSSQALQLPYDPEMEEDSDYSGEPSYPEVFPPLTGYGPELEEEERGV 360

QY 361 KLGLGDFIFYSVLVGKAAATGSDWNTTILACFVAILIGLCTLLLLAVFKKALPALPISI 420
DB 361 KLGLGDFIFYSVLVGKAAATGSDWNTTILACFVAILIGLCTLLLLAVFKKALPALPISI 420

QY 421 TFGLIIFYSTDLNLRPFMDTLASHOLYI 448
DB 421 TFGLIIFYSTDLNLRPFMDTLASHOLYI 448

RESULT 7
W28508
ID W28508 standard; Protein: 447 AA.
AC W28508;
DE 07-DEC-1997 (first entry)
DE Full AD4/AD3LP sequence.
KW AD3; AD4/AD3LP; Alzheimer's disease; chromosome; missegregation;
KW presenilin; inhibitor; AD; trisomy 21; ss.
OS Homo sapiens.
PN W09707213-A2.
PN 27-FEB-1997.
PF 15-AUG-1996; U13314.
PR 16-AUG-1995; US-002448.
PA (HARD) HARVARD COLLEGE.
PI Li J, Potter H;
DR WPI: 97-165297/15.
DR N-PSDB: T87426.
PT Identifying genes which cause chromosome missegregation - useful for
PT identifying causes of and treatments for diseases, e.g. Alzheimer's
PT disease, cancer and ageing
PS Claim 29; Fig 29; 77pp; English.
CC Identifying genes which cause improper chromosome segregation,
CC screening for inhibitors of chromosome missegregation and processes

CC caused by genes encoding chromosome missegregation promoters
CC was exemplified using Alzheimer's disease. The sequences
CC given in T87401 to T87426 can be used in the above methods.
CC It is not clear from the figure legend, the figure and the
CC disclosure of the specification which sequence of Fig 1 and Fig 28
CC is the AD4/AD3LP or the AD3 sequence.
SQ Sequence 447 AA;

Query Match 99.3%; Score 2320.5; DB 1; Length 447;
Best Local Similarity 99.8%; Pred. No. 3.1e-235;
Matches 447; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 MLTFMADSEEEVCDERTSLMSAESPTRSCQEGRGQPEDGENTAQRSENEDEEDP 60
DB 1 MLTFMADSEEEVCDERTSLMSAESPTRSCQEGRGQPEDGENTAQRSENEDEEDP 60

QY 61 DRYVCSGVPGRPPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120
DB 61 DRYVCSGVPGRPPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120

QY 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSMLL 180
DB 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLIMSSMLL 180

QY 181 FLFTYIYLGVLKTYNVAMDYPTLLTWNFGAVGMVCIHWKGPLVLOQAYLIMISALMA 240
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QY 241 LVFTKYLPEWSAWILGAISSYDVLVAVLCPKGLRMLVETAQERNEPIFPALIISSAMVW 300
DB 241 LVFTKYLPEWSAWILGAISSYDVLVAVLCPKGLRMLVETAQERNEPIFPALIISSAMVW 300

QY 301 TVGMAKLDPSSQALQLPYDPEMEEDSDYSGEPSYPEVFPPLTGYGPELEEEERGV 360
DB 301 TVGMAKLDPSSQALQLPYDPEMEEDSDYSGEPSYPEVFPPLTGYGPELEEEERGV 360

QY 361 KLGLGDFIFYSVLVGKAAATGSDWNTTILACFVAILIGLCTLLLLAVFKKALPALPISI 420
DB 361 KLGLGDFIFYSVLVGKAAATGSDWNTTILACFVAILIGLCTLLLLAVFKKALPALPISI 420

QY 421 TFGLIIFYSTDLNLRPFMDTLASHOLYI 448
DB 421 TFGLIIFYSTDLNLRPFMDTLASHOLYI 447

RESULT 8
W05766
ID W05766 standard; Protein: 414 AA.
AC W05766;
DE 25-JUL-1997 (first entry)
DE Presenilin-2 delta263-296 mutation.
DE Presenilin-2; human; hps1-1; hps1-2; ps-2; integral membrane protein; AD;
KW familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;
KW depression; antibody; gene expression modulator; therapy; mutein.
OS Homo sapiens.
FH Key
FT Location/Qualifiers
FT misc_difference 263..264
FT /note= "site of 34 residue deletion"
PN W09634099-A2.
PD 31-OCT-1996.
PF 29-APR-1996; CA0263.
PR 28-APR-1995; US-431048.
PR 28-JUN-1995; US-496841.
PR 31-JUL-1995; US-509359.
PA (HSCR-) HSC RES & DEV LP.
PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
PI Fraser PE, Rommens JM, St George-Hyslop PH;
PI WPI: 96-497631/49.
PT New presenilin genes - useful for diagnosis, therapy and drug
PT screening of familial Alzheimer's disease, cerebral disorders, etc.
PS Claim 4; Page -; 178pp; English.
CC W05763-W05766 represent mutated versions of the human presenilin-2

CC coding sequence for wild type human PS-2. The presenilins are a family of
CC highly conserved integral membrane proteins with a common structural
CC motif, common alternate splicing patterns, and common mutational hot spot
CC regions. Mutations in PS genes are implicated in familial Alzheimer's
CC disease (AD) and possibly other diseases such as cerebral haemorrhage,
CC schizophrenia, depression etc., so detection of mutations in the DNA
CC encoding these sequences can be used for diagnosis of these diseases.
CC These proteins, or vectors that express them or containing antisense
CC sequences, antibodies selective for mutant forms of these proteins (such
CC as W05736) and modulators of PS gene expression are potentially useful
CC for treatment of AD etc. Transgenic animals are useful as models for drug
CC screening. The antibodies can also be used e.g. for affinity purification
CC and in immunoassays.
SQ Sequence 467 AA;

Query Match 62.8%; Score 1468; DB 1; Length 467;
Best Local Similarity 64.0%; Pred. No. 9.9e-146;
Matches 299; Conservative 46; Mismatches 78; Indels 44; Gaps 6;
QY 24 ESPTPRSCQEGRGQEDGENTAQWRSQNEEDGEDDPDRYVCSGVP----GRPPG----- 75
Db 3 EIPAPLSTVFQNMSEDSSAIRSQNSDQERQOQHQRLNDPPIGNGRPQSNRQV 62
QY 75 -----LEELTKYGAHVIMLFPVPTLCMIWVYVATIKSVFYTEKNGOLIYTPTEDT 128
Db 63 VQDSEDEELTKYGAHVIMLFPVPTLCMIWVYVATIKSVFYTEKNGOLIYTPTEDT 122
QY 129 PSVQRLNSVLTLMISVIVVITFLVLYKYRCYKFIHGLWIMSLMLFLFYIYL 188
Db 123 ETVGORALHSILNAAIMISVIVITLLVLYKYRCYKFIHGLWIMSLMLFLFYIYL 182
QY 189 GEVLKTYNVADYPTLLTWNFGAVGVCIIHWKGPLVLOQAYLIMISALMALVFKIYLP 248
Db 183 GEVFTKTYNVADYPTLLTWNFGAVGVCIIHWKGPLVLOQAYLIMISALMALVFKIYLP 242
QY 249 ENSAWVILGAISYDVLVAVLCPKGLRMLVETAQERNEPIFALYISSAMVTVGNKALD 308
Db 243 ENTAWLILAVISYDVLVAVLCPKGLRMLVETAQERNEPIFALYISSAMVTVGNKALD 302
QY 309 PSSQALQLYDPEME-----EDSYDSFGEPSPYVEFPEPLTGYPG----- 350
Db 303 PEAQ--RRVPKPKYNTQARAETQDSGSGNDGFGSEWEAQRDHGLPHRSTPESRAA 360
QY 350 -EEL-----DEEERGVKLGIDPIFYSLVGRAAATGSGDWNNTLACFVAILGLCL 401
Db 361 VOELSGSILTSDPEERGKVLGIDPIFYSLVGRASATASGDWNNTLACFVAILGLCL 420
QY 402 TLLLAVFKKALPALPISITFTGLIFVFTDNLVRFPMDFLASHQLYI 448
Db 421 TLLLAIFKKALPALPISITFTGLVFFVATDYLVPFMDQLAFHQFYI 467

RESULT 11
W23966
ID W23966 standard; Protein: 467 AA.
AC W23966;
DE Mouse presenilin-1 homologue.
KW Presenilin-1; Psl gene; mouse; familial Alzheimer's disease; FAD;
KW cerebral haemorrhage; schizophrenia; depression; epilepsy;
KW mental retardation; diagnosis; therapy; transgenic animal.
OS Mus musculus.
FH Key Location/Qualifiers
FT 82..100
FT Domain /label= TM1
FT /note= "transmembrane domain 1"
FT 101..132
FT Domain /label= TM1-2
FT /note= "hydrophilic loop"
FT 133..154
FT Domain /label= TM2
FT /note= "transmembrane domain 2"

FT Domain 155..163
FT /label= TM2-3
FT /note= "hydrophilic loop"
FT Domain 164..183
FT /label= TM3
FT /note= "transmembrane domain 3"
FT Domain 184..194
FT /label= TM3-4
FT /note= "hydrophilic loop"
FT Domain 195..212
FT /label= TM4
FT /note= "transmembrane domain 4"
FT Domain 213..220
FT /label= TM4-5
FT /note= "hydrophilic loop"
FT Domain 221..238
FT /label= TM5
FT /note= "transmembrane domain 5"
FT Domain 239..243
FT /label= TM5-6
FT /note= "hydrophilic loop"
FT Domain 244..262
FT /label= TM6
FT /note= "transmembrane domain 6"
FT Domain 263..407
FT /label= TM6-7
FT /note= "hydrophilic loop"
FT Domain 408..428
FT /label= TM8
FT /note= "transmembrane domain 8"
FT Misc_difference 177 /note= "Phe177Ser mutation site (Claim 1)"
FT Misc_difference 439 /note= "Ile439Val mutation site (Claim 1)"
PN W09801549-A2.
PD 15-JAN-1998.
PF 04-JUL-1997; CA0475.
PR 02-JAN-1997; US-034590.
PR 05-JUL-1996; US-021673.
PR 12-JUL-1996; US-021700.
PR 08-NOV-1996; US-029895.
PA (HSCR-) HSC RES & DEV LP.
PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
PI Fraser PE, Rommens JW, St George-Hyslop PH;
DR WPI; 98-286355/25.
PT New isolated mutant presenilin-1 genes - useful for developing
PT products for use in detection, diagnosis and therapy of Alzheimer's
PT disease and for drug screening
PS Disclosure; Page 199-200; 238pp; English.
CC This polypeptide comprises the murine presenilin-1 (PS1) homologue.
CC Its amino acid sequence was deduced from an isolated cDNA clone
CC (see W04668). Mutations in the human PS1 and PS2 genes (see
CC W04666-68) have been linked to the development in humans of forms
CC of familial Alzheimer's disease (FAD). All amino acids that are
CC mutated in analysed FAD pedigrees (see W23964) were conserved in
CC the murine homologue. Use of the nucleic acids and proteins
CC comprising or derived from presenilins can be made in screening and
CC diagnosing FAD, identifying and developing therapeutics for
CC treatment of FAD, and in producing cell lines and transgenic
CC animals useful as models of FAD. Methods for identifying
CC substances that bind to, or modulate the activity of a presenilin
CC protein, and methods for identifying substances that affect the
CC interaction of a presenilin-interacting protein with a presenilin
CC protein are also disclosed.
SQ Sequence 467 AA;

Query Match 62.8%; Score 1468; DB 1; Length 467;
Best Local Similarity 64.0%; Pred. No. 9.9e-146;
Matches 299; Conservative 46; Mismatches 78; Indels 44; Gaps 6;
QY 24 ESPTPRSCQEGRGQEDGENTAQWRSQNEEDGEDDPDRYVCSGVP-----GRPPG----- 75

Matches 305; Conservative 40; Mismatches 79; Indels 42; Gaps 8;

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QY 24 ESPTPRSCQGGPGDGTACWRSQENEDGEDPDYVCSGVP-----GRPPG---- 75
Db 3 ELAPLSYFQNAQMSDNHLSNVRSDNRRERQEHDRR-SLGHPEPLSNGRPGNSRQ 61
QY 75 -----LEELTLKYGAKHVIMLFVPTLCMIVVATIKSVREYTKNGQLIYPTFTED 127
Db 62 VWEQDEDEDELTKYGAHVIMLFVPTLCMIVVATIKSVSYTRKDGOLIYPTFTED 121
QY 128 TPSVGBQLNSVNTLIMISIVVMTIFVLVLYKYRCYKFIHGWLMSSMLLFLFYIY 187
Db 122 TETVGRALHSILNAAIMISIVVMTILLVLYKYRCYKVIHAWLIISLLEFFSFIY 181
QY 188 LGEVLKTYNYAMDYPTLLLVWNFAGVWCIIHWKGPLVLQQAAYLIMISALMALVFIKYL 247
Db 182 LGEVFTYNVAVDIITVALLIWNFGVVGMSIHWKGPLRLQQAAYLIMISALMALVFIKYL 241
QY 248 PEWSAWILGAISYDILVAVLCPKPLMLVETAQERNEPIFPALIIYSSAMVTVGMAKL 307
Db 242 PENTAWILLAVISYDILVAVLCPKPLMLVETAQERNEILFPAVIYSSAMVTVGMNAEG 301
QY 308 DPSSQAL--QLPYDPE-MEEDSYDSFGE---PSYPEVFEPPLTGPY----- 350
Db 302 DPEAQRVSKNSKYNASTERESQDTVAENDDGGFSEWEAQRDHLGPHRSTPESRAAV 361
QY 350 EEL-----EEEEGVKLGDFIYSVLVKAATGSDWNTTACFVAILIGLCLT 402
Db 362 QELSSSILAGEDPEERGKLGDFIYSVLVKAATGSDWNTTACFVAILIGLCLT 421
QY 403 LLLIAVFKKALPALPISITFGLIFYSTDLVRPFMDTLASHOLYI 448
Db 422 LLLIAVFKKALPALPISITFGLVYFATDYLQPFMDQLAFHQFYI 467
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Search completed: March 20, 2000, 05:31:20
Job time: 4209 sec

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OM protein - protein search, using sw model

Run on: March 18, 2000, 19:01:42 ; Search time 26.47 Seconds
(without alignments)
225.059 Million cell updates/sec

Title: US-08-509-359B-137
Perfect score: 2336
Sequence: 1 MLTFMADSEEVCDERTSL.....STDNLVRPFMDTLASHQLYI 448

Scoring table: BLOSUM62

Searched: 133990 seqs, 13297546 residues

Database : Issued_Patents_AA.*

Word size : 0

Number of hits that pass the threshold : 133990
1: /cgn2.6/ptodata/2/iaa/5A_COMB.pep.*
2: /cgn2.6/ptodata/2/iaa/5B_COMB.pep.*
3: /cgn2.6/ptodata/2/iaa/6_COMB.pep.*
4: /cgn2.6/ptodata/2/iaa/PCTUS9_COMB.pep.*
5: /cgn2.6/ptodata/2/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	2336	100.0	448	2	US-08-967-101-137
2	2336	100.0	448	2	US-08-592-541-137
3	2320.5	99.3	447	2	US-08-875-972-29
4	1923	82.3	372	2	US-08-967-101-138
5	1923	82.3	372	2	US-08-592-541-138
6	1623.5	69.5	376	2	US-08-875-972-2
7	1457	62.8	467	2	US-08-967-101-134
8	1467	62.8	467	2	US-08-592-541-134
9	1467	62.5	467	3	US-08-670-964-2
10	1461	62.5	467	2	US-08-967-101-2
11	1461	62.5	467	2	US-08-592-541-2
12	1454	62.2	463	3	US-08-670-964-4
13	1438	61.6	407	2	US-08-875-972-4
14	1437	61.5	463	2	US-08-670-479-18
15	1418	60.7	467	2	US-08-367-101-4
16	1418	60.7	467	2	US-08-592-541-4
17	1150	49.2	541	2	US-08-967-101-166
18	1150	49.2	541	2	US-08-592-541-166
19	110	4.7	1294	2	US-08-819-288-3
20	108	4.6	1321	1	US-08-261-822A-3
21	108	4.6	1321	4	PCT-US95-07744A-3
22	101	4.3	1334	2	US-08-986-545-2
23	100	4.3	400	1	US-08-602-010A-8
24	100	4.3	400	1	US-08-680-726A-8
25	95	4.1	391	1	US-07-816-283-4
26	95	4.1	391	1	US-08-417-103-4
27	93	4.0	391	1	US-07-816-283-2
28	93	4.0	391	1	US-08-417-103-2
29	93	4.0	391	1	US-08-417-103-14
30	91	3.9	509	2	US-08-331-392-6
31	88.5	3.8	452	1	US-08-117-361C-1
32	88.5	3.8	3169	2	US-08-477-451-6
33	87	3.7	492	2	US-08-355-844-3
34	87	3.7	492	4	PCT-US95-16126-3

35 86.5 3.7 1480 1 US-07-637-621-2 Sequence 2, Appli
36 86.5 3.7 1480 1 US-08-136-742A-2 Sequence 2, Appli
37 86.5 3.7 1480 1 US-08-135-809A-2 Sequence 2, Appli
38 86.5 3.7 1480 2 US-08-951-912-2 Sequence 2, Appli
39 86.5 3.7 1479 2 US-08-951-912-4 Sequence 4, Appli
40 86.5 3.7 1480 2 US-08-951-912-6 Sequence 6, Appli
41 86.5 3.7 1480 2 US-08-691-605-2 Sequence 2, Appli
42 86.5 3.7 1480 2 US-08-455-552A-14 Sequence 14, Appli
43 86.5 3.7 1480 4 PCT-US93-11667-2 Sequence 2, Appli
44 86.5 3.7 1480 5 5240846-5 Patent No. 5240846
45 86 3.7 413 2 US-08-808-793-25 Sequence 25, Appli

ALIGNMENTS

RESULT 1
US-08-967-101-137
; Sequence 137, Application US/08967101
; Patent No. 5840540
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/967,101
; FILING DATE: 10-NOV-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/592,541
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 137:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 448 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-967-101-137

Query Match 100.0%; Score 2336; DB 2; Length 448;
Best Local Similarity 100.0%; Pred. No. 1e-231;
Matches 448; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MLTFMADSEEVCDERTSLMSAESPTPRSCQGRQGPEDGNTAQWRSQNEEDGEDP 60
Db 1 MLTFMADSEEVCDERTSLMSAESPTPRSCQGRQGPEDGNTAQWRSQNEEDGEDP 60
QY 61 DRYVCSGVPGPPGLEELTLKYGAKHVMFLPVPVTLCMIVVVATIKSVRYTERNGQLI 120
Db 61 DRYVCSGVPGPPGLEELTLKYGAKHVMFLPVPVTLCMIVVVATIKSVRYTERNGQLI 120

US-08-875-972-29

Query Match 99.3%; Score 2320.5; DB 2; Length 447;
Best Local Similarity 99.8%; Pred. No. 4e-230;
Matches 447; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 MLTPMADSEEEVCDERTSLMSAESPTRSCQGRQGPEDGENTAQRNSQNEEDGEDP 60
DB 1 MLTPMADSEEEVCDERTSLMSAESPTRSCQGRQGPEDGENTAQRNSQNEEDGEDP 60
QY 61 DRYVCSGVPGRPGLEELIKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120
DB 61 DRYVCSGVPGRPGLEELIKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLI 120
QY 121 YTPETEDPSVQGRLLNSVNTLMISVIVVMTIFLVLYKYRCYKFTHGWLMSLML 180
DB 121 YTPETEDPSVQGRLLNSVNTLMISVIVVMTIFLVLYKYRCYKFTHGWLMSLML 180
QY 181 FLFTYIYLGVLKTYNVAMDYPTLLTWNFAGVMCIHWKGPLVLOQAYLIMISALMA 240
DB 181 FLFTYIYLGVLKTYNVAMDYPTLLTWNFAGVMCIHWKGPLVLOQAYLIMISALMA 240
QY 241 LVFTKYLPEWSAWVILGAISYIDLAVLCPRGLMVLVETAQERNEPFPALIIYSSAMVW 300
DB 241 LVFTKYLPEWSAWVILGAISYIDLAVLCPRGLMVLVETAQERNEPFPALIIYSSAMVW 300
QY 301 TVGMKLDPSOGALQLPYDPEMEDSDYSFGPSYFEPFPLTGYPGSELEEEERG 360
DB 301 TVGMKLDPSOGALQLPYDPEMEDSDYSFGPSYFEPFPLTGYPGSELEEEERG 360
QY 361 KLGLGDFIFYSVLVGRKAAATGSGDWNNTLACFVAILGLCTLALLAVFKKALPALPISI 420
DB 360 KLGLGDFIFYSVLVGRKAAATGSGDWNNTLACFVAILGLCTLALLAVFKKALPALPISI 420
QY 421 TFLGFIYFSTDNLVRPMDTLASHOLYI 448
DB 420 TFLGFIYFSTDNLVRPMDTLASHOLYI 447

RESULT 4

US-08-967-101-138
; Sequence 138, Application US/08967101
; Patent No. 5840540
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/967,101
; FILING DATE: 10-NOV-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/592,541
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:

TELEPHONE: (617) 248-7000

TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 138:

SEQUENCE CHARACTERISTICS:

LENGTH: 372 amino acids

TYPE: amino acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: protein

US-08-967-101-138

Query Match 82.3%; Score 1923; DB 2; Length 372;

Best Local Similarity 100.0%; Pred. No. 2e-189;

Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 77 BELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLIYTPFTEDTPSVQORLL 136
DB 1 BELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGOLIYTPFTEDTPSVQORLL 60
QY 137 NSVLTLMISVIVVMTIFLVLYKYRCYKFTHGWLMSLMLFLTYYILGEVLKTYN 196
DB 61 NSVLTLMISVIVVMTIFLVLYKYRCYKFTHGWLMSLMLFLTYYILGEVLKTYN 120
QY 197 VAMDYPTLLTWNFAGVMCIHWKGPLVLOQAYLIMISALMALVFIKYLPWSAWVIL 256
DB 121 VAMDYPTLLTWNFAGVMCIHWKGPLVLOQAYLIMISALMALVFIKYLPWSAWVIL 180
QY 257 GAISYIDLAVLCPRGLMVLVETAQERNEPFPALIIYSSAMVWTVGMKLDPSOGALQ 316
DB 181 GAISYIDLAVLCPRGLMVLVETAQERNEPFPALIIYSSAMVWTVGMKLDPSOGALQ 240
QY 317 LPYDPEMEDSDYSFGPSYFEPFPLTGYPGSELEEEERGKLGDFIFYSVLVGR 376
DB 241 LPYDPEMEDSDYSFGPSYFEPFPLTGYPGSELEEEERGKLGDFIFYSVLVGR 300
QY 377 AAATGSGDWNNTLACFVAILGLCTLALLAVFKKALPALPISITFGLIFYSTDLNVRP 436
DB 301 AAATGSGDWNNTLACFVAILGLCTLALLAVFKKALPALPISITFGLIFYSTDLNVRP 360
QY 437 FMDTLASHOLYI 448
DB 361 FMDTLASHOLYI 372

RESULT 5

US-08-592-541-138
; Sequence 138, Application US/08592541
; Patent No. 5986054
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
; STREET: High Street Tower - 125 High Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/592,541
; FILING DATE:
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:

SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/967,101
FILING DATE: 10-NOV-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/592,541
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Pitcher, Edmund R.
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ ID NO: 134:
SEQUENCE CHARACTERISTICS:
LENGTH: 467 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-967-101-134

Query Match 62.8%; Score 1467; DB 2; Length 467;
Best Local Similarity 65.7%; Pred. No. 2e-142;
Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;

QY 24 ESPTRSCQGRGQPEDGNTAQWRSQNEDEEDDPDRYVCSGVP-----GRPPG---- 75
Db 3 ELPAFLSYFQNAQMSDNHLSNTRVSDNRRERQEHDRR-SLGHPEPLSNRQGNRSRQ 61
QY 75 -----LEELTLKYGAKHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYPTFTED 127
Db 62 VVEQDEEDELTLKYGAKHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYPTFTED 121
QY 128 TPSVGORLNSVLNTLMISVIVMTIFLVLYKYRCYKFIHGLWLMSSLMILFLFTYIY 187
Db 122 TETVGORALHSILNAAIMSVIVMTIFLVLYKYRCYKFIHGLWLMSSLMILFLFTYIY 181
QY 188 LGEVLKTYNVAMDYPTLLTVMNFGAVGVCIHMKGPLVQLQAYLIMISALMALVFIKYL 247
Db 182 LGEVFTYNAVVDYITVALLIWNFGVGMISIHMKGPLRQLQAYLIMISALMALVFIKYL 241
QY 248 PWSAWILGALSVDLVAVLCPKPLRMVETAQERNPIFIPALYSSAMVTVGMAKL 307
Db 242 PWTAWLILAVISVDLVAVLCPKPLRMVETAQERNPIFIPALYSSAMVTVGMAKL 301
QY 308 DPSSQCAL--QLPYDPE-MEEDSYDSFGE---PSYPEVFPPLTGYPG----- 350
Db 302 DPEAQRVSKNSKYNASTERSQDTVAENDDGGFSEWEAQRDHLGPHRSTPESRAAV 361
QY 350 EEL-----EEEEERGVKLGDFIFYSVLVGAATGSDWNTTACFVAILIGLCLT 402
Db 362 QELSSSILAGEDPEERGVKLGDFIFYSVLVGAATGSDWNTTACFVAILIGLCLT 421
QY 403 LLLAVFKKALPALPISITIFGLIFYSTDLNLRPFMDTLASHQLYI 448
Db 422 LLLAIFKKALPALPISITIFGLVYFATDYLQVPMQDLAFHQFYI 467

RESULT 8
US-08-592-541-134
Sequence 134, Application US/08592541
Patent No. 5986054
GENERAL INFORMATION:
APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: BOWMENS, JOHANNA M
APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183
CORRESPONDENCE ADDRESS:
ADDRESSEE: TESTA, HURWITZ & THIBEAULT

STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/592,541
FILING DATE:
CLASSIFICATION: 800
ATTORNEY/AGENT INFORMATION:
NAME: Pitcher, Edmund R.
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ ID NO: 134:
SEQUENCE CHARACTERISTICS:
LENGTH: 467 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-592-541-134

Query Match 62.8%; Score 1467; DB 2; Length 467;
Best Local Similarity 65.7%; Pred. No. 2e-142;
Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;

QY 24 ESPTRSCQGRGQPEDGNTAQWRSQNEDEEDDPDRYVCSGVP-----GRPPG---- 75
Db 3 ELPAFLSYFQNAQMSDNHLSNTRVSDNRRERQEHDRR-SLGHPEPLSNRQGNRSRQ 61
QY 75 -----LEELTLKYGAKHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYPTFTED 127
Db 62 VVEQDEEDELTLKYGAKHVIMLFVPTLCMIVVATIKSVRYTEKNGQLIYPTFTED 121
QY 128 TPSVGORLNSVLNTLMISVIVMTIFLVLYKYRCYKFIHGLWLMSSLMILFLFTYIY 187
Db 122 TETVGORALHSILNAAIMSVIVMTIFLVLYKYRCYKFIHGLWLMSSLMILFLFTYIY 181
QY 188 LGEVLKTYNVAMDYPTLLTVMNFGAVGVCIHMKGPLVQLQAYLIMISALMALVFIKYL 247
Db 182 LGEVFTYNAVVDYITVALLIWNFGVGMISIHMKGPLRQLQAYLIMISALMALVFIKYL 241
QY 248 PWSAWILGALSVDLVAVLCPKPLRMVETAQERNPIFIPALYSSAMVTVGMAKL 307
Db 242 PWTAWLILAVISVDLVAVLCPKPLRMVETAQERNPIFIPALYSSAMVTVGMAKL 301
QY 308 DPSSQCAL--QLPYDPE-MEEDSYDSFGE---PSYPEVFPPLTGYPG----- 350
Db 302 DPEAQRVSKNSKYNASTERSQDTVAENDDGGFSEWEAQRDHLGPHRSTPESRAAV 361
QY 350 EEL-----EEEEERGVKLGDFIFYSVLVGAATGSDWNTTACFVAILIGLCLT 402
Db 362 QELSSSILAGEDPEERGVKLGDFIFYSVLVGAATGSDWNTTACFVAILIGLCLT 421
QY 403 LLLAVFKKALPALPISITIFGLIFYSTDLNLRPFMDTLASHQLYI 448
Db 422 LLLAIFKKALPALPISITIFGLVYFATDYLQVPMQDLAFHQFYI 467

RESULT 9
US-08-670-964-2
Sequence 2, Application US/08670964
Patent No. 6010874
GENERAL INFORMATION:
APPLICANT: Hardy, John A.
TITLE OF INVENTION: EARLY ONSET ALZHEIMER'S DISEASE

```

Query Match      62.5%; Score 1461; DB 2; Length 467;
Best Local Similarity 65.5%; Pred. No. 8.1e-142;
Matches 305; Conservative 39; Mismatches 80; Indels 42; Gaps 8;

QY      24  ESTPTSCQGGROGPEGNTAQRWSENEDEGDEDPDHYCVSGVP-----GRRPG---- 75
      | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db      3  ELFAPLSYFQNAQMSDENHLSNTRVRSQNDNREGRHEHNDRR-SLGHPEPLSNRGRCQNSRQ 61
      | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY      75  -----LEBELTKYGAKHVIMLFVPVTLTCMIVVYVATIKSVREYTEKNGOLITYPTFED 127
      : | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db      62  VWEQDEEDEBELTKYGAKHVIMLFVPVTLTCMIVVYVATIKSVREYTRKDGQLITYPTFED 121
      | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY      128  TPSVQGRLLNSVLNTLIMISIVVYVMTFLVLYLYKRCYKFTGHGLIMSSLMMLFLFYIY 187
      | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db      122  TETVQGRALSHLSNAALIMISIVVYVMTILLVYLYKRCYKVTHAWLIMLSLLLLLFFSFYI 181
      | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY      188  LGEVLKTYNYAMDPTLLLTWNFGVGMVCIHWKGPVLVQQAYLIMISALMALVFIKYL 247

```

Query Match 62.2%; Score 1454; DB 3; Length 463;
Best Local Similarity 55.3%; Pred. No. 4.2e-141;
Matches 303; Conservative 40; Mismatches 79; Indels 42; Gaps 8;
QY 24 ESTPRSCGROGPEGNTAOWRSENEEDGEDPDYVCSGVP---GRPPG----- 75

REFERENCE/DOCKET NUMBER: P50361
TELEPHONE: 610-270-5219
TELEFAX: 610-270-5090
TELEX:

INFORMATION FOR SEQ ID NO: 18:

SEQUENCE CHARACTERISTICS:

LENGTH: 463 amino acids

TYPE: amino acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: peptide

HYPOTHETICAL: NO

ANTI-SENSE: NO

FRAGMENT TYPE: N-terminal

ORIGINAL SOURCE:

US-08-670-479-18

Query Match 61.5%; Score 1437; DB 2; Length 463;
Best Local Similarity 64.9%; Pred. No. 2.3e-139;
Matches 301; Conservative 41; Mismatches 80; Indels 42; Gaps 8;

QY 24 ESPTPRSCQGRQGEDGENTAQRSENEDEEDDPDYVCSGVP---GRPPG----- 75
DB 3 ELPAPLSYFQNAQMSDNHLS---NTNDRERQEHNDRLSGHPPEPLSNRGPQNSRQV 59
QY 75 -----LEELTKYGAHVIMLVFVPTLCMVVATIKSVFYTEKNGOLIYPTPTEDTP 129
DB 60 EQDEEDELTKYGAHVIMLVFVPTLCMVVATIKSVFYTRKDGOLIYPTPTEDTE 119
QY 130 SVGORLNSVNTLMISVIVMTILVLYKYRKYKIHWGLMSSLLMLFLFYIYL 189
DB 120 TVGORLNSVNTLMISVIVMTILVLYKYRKYKIHWGLMSSLLMLFLFYIYL 179
QY 190 EVLKTYNVAMDYPTLLLVFNWFGAVGVCIIHWKGLPLVQQAAYLIMISALMALVFIKYLPE 249
DB 180 EVLKTYNVAMDYPTLLLVFNWFGAVGVCIIHWKGLPLVQQAAYLIMISALMALVFIKYLPE 239
QY 250 WSAWVILGASVYDLVAVLCPKGLRMLVETAQRNEPIFFPALIYSSAMVTVGMKLD 309
DB 240 WTAWLILAVISYDLVAVLCPKGLRMLVETAQRNEPIFFPALIYSSAMVTVGMKLD 299
QY 310 SSQAL--QLPYDPE--MEEDSYDFGE--PSYPVFPEPLTGYPG-----EE 351
DB 300 EAQRVSKNSYNAESTRESQDTVAENDDGGFSEWEAQRDLSHLGPHRSTPESRAAVQE 359
QY 352 L-----EEEEERGVKLGIDGFIFYSVLVGKAAATGSDWNTTLACFVAILIGLCLTLL 404
DB 360 LSSSLAGEDPEERGVKLGIDGFIFYSVLVGKAAATGSDWNTTLACFVAILIGLCLTLL 419
QY 405 LLAVFKKALPALISITFGLIFYSTDLNVRPMDTLASHQLYI 448
DB 420 LLAVFKKALPALISITFGLIFYSTDLNVRPMDTLASHQLYI 463

RESULT 15

US-08-967-101-4

Sequence 4, Application US/08967101

Patent No. 5840540

GENERAL INFORMATION:

APPLICANT: ST. GEORGE-HYSLOP, PETER H

APPLICANT: ROMMENS, JOHANNA M

APPLICANT: FRASER, PAUL E

TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED

TITLE OF INVENTION: TO ALZHEIMER'S DISEASE

NUMBER OF SEQUENCES: 183

CORRESPONDENCE ADDRESS:

ADDRESSEE: TESTA, HURWITZ & THIBEAULT

STREET: High Street Tower - 125 High Street

CITY: Boston

STATE: Massachusetts

COUNTRY: U.S.A.

ZIP: 02110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/967,101
FILING DATE: 10-NOV-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/592,541
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Pitcher, Edmund R.
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 467 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-967-101-4

Query Match 60.7%; Score 1418; DB 2; Length 467;
Best Local Similarity 61.9%; Pred. No. 2.1e-137;
Matches 289; Conservative 48; Mismatches 86; Indels 44; Gaps 6;

QY 24 ESPTPRSCQGRQGEDGENTAQRSENEDEEDDPDYVCSGVP---GRPPG----- 75
DB 3 ELPAPLSYFQNAQMSDNHLS---NTNDRERQEHNDRLSGHPPEPLSNRGPQNSRQV 62
QY 75 -----LEELTKYGAHVIMLVFVPTLCMVVATIKSVFYTEKNGOLIYPTPTEDTP 128
DB 63 EQDEEDELTKYGAHVIMLVFVPTLCMVVATIKSVFYTRKDGOLIYPTPTEDT 122
QY 129 PSVGORLNSVNTLMISVIVMTILVLYKYRKYKIHWGLMSSLLMLFLFYIYL 188
DB 123 TVGORLNSVNTLMISVIVMTILVLYKYRKYKIHWGLMSSLLMLFLFYIYL 182
QY 189 EVLKTYNVAMDYPTLLLVFNWFGAVGVCIIHWKGLPLVQQAAYLIMISALMALVFIKYL 248
DB 183 EVLKTYNVAMDYPTLLLVFNWFGAVGVCIIHWKGLPLVQQAAYLIMISALMALVFIKYL 242
QY 249 WSAWVILGASVYDLVAVLCPKGLRMLVETAQRNEPIFFPALIYSSAMVTVGMKLD 308
DB 243 WTAWLILAVISYDLVAVLCPKGLRMLVETAQRNEPIFFPALIYSSAMVTVGMKLD 302
QY 309 PSQALQLPYDPE-----EDSYDFGEPSPYVPEPPLTGYPG----- 350
DB 303 PEAQ--RRVPKPKYNTQARERQSDSGNDGSEWEAQRDLSHLGPHRSTPESRAA 360
QY 350 -EEL-----EEEEERGVKLGIDGFIFYSVLVGKAAATGSDWNTTLACFVAILIGLCL 401
DB 361 VQELSSILTSDEPERGVKLGIDGFIFYSVLVGKAAATGSDWNTTLACFVAILIGLCL 420
QY 402 TLLLVAVFKKALPALISITFGLIFYSTDLNVRPMDTLASHQLYI 448
DB 421 XLLLLAIYKKGXPAPISITFTGFGVFAFDYLVQPMQDLAFHQFYI 467

Search completed: March 18, 2000, 19:55:31
Job time: 3229 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 14:03:43 ; Search time 41.25 Seconds
(without alignments)
512.287 Million cell updates/sec

Title: US-08-509-359B-137
Perfect score: 2336
Sequence: 1 MLTFWASDSEEEVCDERTSL.....STDNLVRFPMDTLASHQLYI 448

Scoring table: BLOSUM62

Searched: 142080 seqs, 47169319 residues

Database : PIR_62.*

Word size : 0

Number of hits that pass the threshold : 142080

- 1: pir1.*
- 2: pir2.*
- 3: pir3.*
- 4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	2336	100.0	448	2	I58098	E5-1 protein - hum
2	2328	99.7	448	2	A56993	presenilin 2 - hum
3	2215.5	94.8	442	2	I39174	seven trans-membra
4	1624.5	69.5	449	2	JC5391	presenilin-beta -
5	1468	62.8	467	2	I78388	5182 protein - mou
6	1467	62.8	467	2	S58396	presenilin 1, spll
7	1454	62.2	463	2	S63883	presenilin I-463 -
8	1449.5	62.1	433	2	JC5390	presenilin-alpha -
9	1447	61.9	467	2	JC5080	presenilin 1 prote
10	1440	61.6	463	2	JC5081	presenilin 1 prote
11	1035.5	44.3	374	2	S63884	presenilin 1, spll
12	967	41.4	461	2	S60253	sel-12 protein - C
13	524.5	22.5	358	2	T15184	hypothetical prote
14	513	22.0	453	2	T00724	presenilin homolog
15	274	11.7	465	2	A43459	sperm membrane pro
16	119	5.1	531	2	T11596	hypothetical prote
17	110	4.7	2016	2	A38195	sodium channel pro
18	109	4.7	1840	1	CHRTM1	sodium channel pro
19	106	4.5	398	2	H75043	mg2+ transport pro
20	104	4.5	1476	1	A39901	cystic fibrosis tr
21	104	4.5	1476	1	A40303	cystic fibrosis tr
22	104	4.5	826	2	T02268	potassium transpor
23	103.5	4.4	1450	2	JC6139	cystic fibrosis tr
24	103	4.4	382	2	A47882	ubiquinol--cytochr
25	102.5	4.4	893	2	A47550	bride of sevenless
26	102	4.4	379	2	I48135	ubiquinol--cytochr
27	101.5	4.3	1681	2	A55138	sodium channel mRNA
28	100.5	4.3	1951	2	S00320	sodium channel pro
29	100.5	4.3	1983	2	A60054	isp4 protein homol
30	100.5	4.3	766	2	T01900	integrin-associate
31	100.5	4.3	324	2	S36646	cytochrome-c oxida
32	100	4.3	592	2	E70488	serotonin receptor
33	99.5	4.3	379	2	JC6178	sodium channel pro
34	99	4.2	2019	2	A33996	probable amino aci
35	99	4.2	521	2	T11710	

36 98 4.2 461 2 T11829 NADH dehydrogenase
37 98 4.2 1836 2 J50648 sodium channel alp
38 98 4.2 1836 2 I51964 sodium channel alp
39 98 4.2 1836 2 I64893 sodium channel alp
40 98 4.2 1835 2 I54323 sodium channel alp
41 98 4.2 217 2 S01095 hypothetical prote
42 98 4.2 1695 2 JE0084 voltage-gated sodi
43 97.5 4.2 441 2 S13425 endothelin recepto
44 97 4.2 381 2 T11440 ubiquinol--cytochr
45 97 4.2 447 2 S52968 NADH dehydrogenase

ALIGNMENTS

RESULT 1

I58098
E5-1 protein - human
C:Species: Homo sapiens (man)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 29-Sep-1999
C:Accession: I58098
R:Rogaev, E.I.; Sherrington, R.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; Liang, Y.; C
.; Cohen, D.; Lannfelt, L.; Fraser, P.E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 376, 775-778, 1995
A:Title: Familial Alzheimer's disease in kindreds with missense mutations in a gene o
A:Reference number: I58098; MUID:95379971
A:Accession: I58098
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: mRNA
A:Residues: 1-448 <RES>
A:Cross-references: GB:I44577; NID:g950347; PIDN:AAC42012.1; PID:g950348
C:Genetics:
A:Gene: E5-1
C:Superfamily: presenilin

Query Match 100.0%; Score 2336; DB 2; Length 448;
Best Local Similarity 100.0%; Pred. No. 8.1e-167;
Matches 448; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 MLTFWASDSEEEVCDERTSLMSAESPTPRSCQEGRGQPEDGENTAQWRSQENDEEDP 60
Db 1 MLTFWASDSEEEVCDERTSLMSAESPTPRSCQEGRGQPEDGENTAQWRSQENDEEDP 60
Qy 61 DRYVCSGVPGRPGLLEELTLKYGAKHVIMLFVPTLCMIVVVVATIKSVRYTEKNGQLI 120
Db 61 DRYVCSGVPGRPGLLEELTLKYGAKHVIMLFVPTLCMIVVVVATIKSVRYTEKNGQLI 120
Qy 121 YPTPTDTPSYGQRLNLSVNTLIMISIVVMTIFLVLYKYRCYKFHGWLIMSLMLL 180
Db 121 YPTPTDTPSYGQRLNLSVNTLIMISIVVMTIFLVLYKYRCYKFHGWLIMSLMLL 180
Qy 181 FLFYIYLGEVLKTYNAMDYPTLLLTWNFGAVGMVCIHKKGPLVLOQAYLIMISALMA 240
Db 181 FLFYIYLGEVLKTYNAMDYPTLLLTWNFGAVGMVCIHKKGPLVLOQAYLIMISALMA 240
Qy 241 LVFIKLPESAWILGSAISYDVLVAVLCPKGPLRMLVETAQERNEPIFPALIISSAMVW 300
Db 241 LVFIKLPESAWILGSAISYDVLVAVLCPKGPLRMLVETAQERNEPIFPALIISSAMVW 300
Qy 301 TVGMAKLDPSQGAQLPYDPMEEDSDSFGESPSYVEVFEPPLTGYGPEEELEEEERG 360
Db 301 TVGMAKLDPSQGAQLPYDPMEEDSDSFGESPSYVEVFEPPLTGYGPEEELEEEERG 360
Qy 361 KLGLGDFIFYSVLGVKAAATGSGDNTTLACFVAILGLICLTLTLLLAVEKKALPALPISI 420
Db 361 KLGLGDFIFYSVLGVKAAATGSGDNTTLACFVAILGLICLTLTLLLAVEKKALPALPISI 420
Qy 421 TFGLIFFYSTNLVRFPMDTLASHQLYI 448
Db 421 TFGLIFFYSTNLVRFPMDTLASHQLYI 448

Qy 7 SDSEEVCDERTSLMSAESPTPRSCQEGRQGEDGENTAQRWSENEEDGEDDPRYVCS 66
Db 5 SDSEDEECNERTSLITSESPPLPSVQDGVQASEGLETSYHREORQSDTQNNED----- 58
Qy 67 GVP-GRPGCL-----EEELTKYGAHVIMLFVPTVLCMIVVATIKSVRYTT 113
Db 58 -VPNGRTSGADAYNSETTVNEEEELTKYGAHVIMLFVPTVLCMIVVATIKSVRYTT 116
Qy 114 EKNGLIYTPETEDTPSYGQRLNLSVNTLMISVIVMTIFLVLYKYKCFHGLI 173
Db 117 EKDGLIYTPESDTPSYGRLNLSVNTLMISVIVMTIFLVLYKYKCFHGLI 176
Qy 174 MSSMLLEFLFYIYLGELVKYNNVAMDYPTLLTWNVFAGVGMVCIHWKGPLVLOQAYLI 233
Db 177 LSSMLLEFLFYIYLGELVKYNNVAMDYPTLLTWNVFAGVGMVCIHWKGPLVLOQAYLI 236
Qy 234 MISALMALVFIKYLPEWSAWVILGAISYDVLAVLCPLKPLRMLVETAQRNEPIFPALI 293
Db 237 MISALMALVFIKYLPEWSAWVILGAISYDVLAVLCPLKPLRMLVETAQRNEPIFPALI 296
Qy 294 YSSAWVTVGMAKLDPSQGL--QLPY-----DPEMEDSDTSFGEPSPYEPFEPPLTGY 347
Db 297 YSSAWVTVGMAKLDPSQGL--QLPY-----DPEMEDSDTSFGEPSPYEPFEPPLTGY 347
Qy 348 PGELEEEERGVKLGIDGFIFYSVLVGKAAATGSGDWNNTTLACFVAILIGLCLTLLLLA 407
Db 349 NLSEDDPEERGVKLGIDGFIFYSVLVGKAAATGSGDWNNTTLACFVAILIGLCLTLLLLA 408
Qy 408 VFKKALPALPISITFGLIFYFSTDNLRPFMDTLASHOLYI 448
Db 409 VFKKALPALPISITFGLIFYFSTDNLRPFMDTLASHOLYI 449
RESULT 5
S182 protein - mouse
C:Species: Mus musculus (house mouse)
C:Date: 27-Feb-1997 #sequence_revision 27-Feb-1997 #text_change 29-Sep-1999
C:Accession: 178388
R:Sherrington, R.; Rogava, E.I.; Liang, Y.; Rogava, E.A.; Levesque, G.; Ikeda, M.; Chiro,
ero, I.; Pinessi, L.; Nee, L.; Chumakov, I.; Pollen, D.; Brookes, A.; Sanseau, P.; Pollen,
E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 375, 754-760, 1995
A:Title: Cloning of a gene bearing missense mutations in early-onset familial Alzheimer
A:Reference number: 158095; MUID:95319502
A:Accession: 178388
A:Molecule type: mRNA
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Residues: 1-467 <RES>
A:Cross-references: GB:LA2177; NID:9904129; PIDN:AAC42094.1; PID:9904130
C:Superfamily: presenilin

Query Match 62.8%; Score 1468; DB 2; Length 467;
Best Local Similarity 64.08; Pred. No. 3.9e-102;
Matches 299; Conservative 46; Mismatches 78; Indels 44; Gaps 6;
Qy 24 ESPTPRSCQEGRQGEDGENTAQRWSENEEDGEDDPRYVCSGVP-----GRPPG----- 75
Db 3 EIPAPLSYFQNAQMSDSSSHSSAIRSQNDSQERQOQDRQLNDPEPISNGRPSQNSRQV 62
Qy 75 -----LEEELTKYGAHVIMLFVPTVLCMIVVATIKSVRYTEKNGQLIYTPFTEDT 128
Db 63 VEQDEEDELTKYGAHVIMLFVPTVLCMIVVATIKSVRYTEKNGQLIYTPFTEDT 122
Qy 129 PSVGORLNSVNTLMISVIVMTIFLVLYKYKCFHGLIYLYL 188
Db 123 ETVGORLNSVNTLMISVIVMTIFLVLYKYKCFHGLIYLYL 182
Qy 189 GEVLTATYVNDYPTLLTWNVFAGVGMVCIHWKGPLVLOQAYLIMISALMALVFIKILP 248
Db 183 GEVLTATYVNDYPTLLTWNVFAGVGMVCIHWKGPLVLOQAYLIMISALMALVFIKILP 242

Qy 249 EWSAWVILGAISYDVLAVLCPLKPLRMLVETAQRNEPIFPALYSSAWVTVGMAKLD 308
Db 243 EWTAWLILAVISYDVLAVLCPLKPLRMLVETAQRNETLFPALYSSAWVTVGMAKLD 302
Qy 309 PSSQALQLPYDPEME-----EDSYDSFGEPSPYEPFEPPLTGYPG----- 350
Db 303 PEAQ--RRVPKPKYNTQRAERETQDSGSGNDGDFSEWEAQRDQSHLGRHSTPESRAA 360
Qy 350 -BEL-----EEERGVKLGIDGFIFYSVLVGKAAATGSGDWNNTTLACFVAILIGLCL 401
Db 361 VOELSGSILTSDEPBERGVKLGIDGFIFYSVLVGKASATASGDWNNTTLACFVAILIGLCL 420
Qy 402 TLLLAVERKALPALPISITFGLIFYFSTDNLRPFMDTLASHOLYI 448
Db 421 TLLLAVERKALPALPISITFGLIFYFSTDNLRPFMDTLASHOLYI 467
RESULT 6
S58396
Presenilin 1, splice form 467 - human
N:Alternate names: Alzheimer's disease protein 3; protein S182
C:Species: Homo sapiens (man)
C:Date: 29-Jan-1998 #sequence_revision 13-Feb-1998 #text_change 29-Sep-1999
C:Accession: S58396; S71401; S71402
R:Sherrington, R.; Rogava, E.I.; Liang, Y.; Rogava, E.A.; Levesque, G.; Ikeda, M.; Chiro,
ero, I.; Pinessi, L.; Nee, L.; Chumakov, I.; Pollen, D.; Brookes, A.; Sanseau, P.; Po
E.; Rommens, J.M.; St George-Hyslop, P.H.
Nature 375, 754-760, 1995
A:Title: Cloning of a gene bearing missense mutations in early-onset familial Alzhelm
A:Reference number: 158095; MUID:95319502
A:Accession: S58396
A:Molecule type: mRNA
A:Residues: 1-467 <SHE>
A:Cross-references: EMBL:LA2110; NID:9904118; PIDN:AAB46416.1; PID:9904119
R:Vidal, R.; Ghiso, J.; Wisniewski, T.; Frangione, B.
FEBS Lett. 393, 19-23, 1996
A:Title: Alzheimer's presenilin 1 gene expression in platelets and megakaryocytes. Id
A:Reference number: S71401; MUID:96397521
A:Accession: S71401
A:Status: not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 24-32;254-256, 290-292;316-317, 376-379 <VIW>
A:Experimental source: Dami megakaryotic cell line (ATCC CRL-9792) and platelets
C:Genetics:
A:Gene: GDB:PSEN1; AD3; PAD; S182; PS1
A:Cross-references: GDB:135682; OMIM:104311
A:Map position: 14q24.3-14q24.3
C:Superfamily: presenilin
C:Keywords: alternative splicing; Alzheimer's disease; glycoprotein; transmembrane pr
F:82-100/Domain: transmembrane #status predicted <TM1>
F:133-154/Domain: transmembrane #status predicted <TM2>
F:164-185/Domain: transmembrane #status predicted <TM3>
F:195-213/Domain: transmembrane #status predicted <TM4>
F:221-238/Domain: transmembrane #status predicted <TM5>
F:244-264/Domain: transmembrane #status predicted <TM6>
F:281-301/Domain: transmembrane #status predicted <TM7>
F:408-428/Domain: transmembrane #status predicted <TM8>
F:433-453/Domain: transmembrane #status predicted <TM9>
F:279,405/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 62.8%; Score 1467; DB 2; Length 467;
Best Local Similarity 65.7%; Pred. No. 4.6e-102;
Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;
Qy 24 ESPTPRSCQEGRQGEDGENTAQRWSENEEDGEDDPRYVCSGVP-----GRPPG----- 75
Db 3 EIPAPLSYFQNAQMSDSSSHSSAIRSQNDSQERQOQDRQLNDPEPISNGRPSQNSRQ 61
Qy 75 -----LEEELTKYGAHVIMLFVPTVLCMIVVATIKSVRYTEKNGQLIYTPFTED 127
Db 62 VEQDEEDELTKYGAHVIMLFVPTVLCMIVVATIKSVRYTEKNGQLIYTPFTED 121

presenilin 1 protein isoform 467 - lesser mouse lemur
C:Species: Microcebus murinus (lesser mouse lemur)
C:Date: 31-Jan-1997 #sequence_revision 31-Jan-1997 #text_change 29-Sep-1999
C:Accession: JC5080
R:Calenda, A.; Mestre-Frances, N.; Czech, C.; Pradier, L.; Petter, A.; Bons, N.; Bellis,
Biochem. Biophys. Res. Commun. 228, 430-439, 1996
A:Title: Molecular cloning, sequencing, and brain expression of the presenilin 1 gene in
A:Reference number: JC5080; MUID:97079199
A:Accession: JC5080
A:Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-467 <CAL>
A:Cross-references: EMBL:Z71333; NID:g1707591; PIDN:CAA95930.1; PID:e248527; PID:g170759
A:Experimental source: brain
C:Comment: This protein is an integral membrane protein with seven transmembrane domain
C:Genetics:
A:Gene: psi
A:Map position: 14
C:Superfamily: presenilin
C:Keywords: transmembrane protein
F:82-100/Domain: transmembrane #status predicted <TM1>
F:133-154/Domain: transmembrane #status predicted <TM2>
F:164-185/Domain: transmembrane #status predicted <TM3>
F:195-213/Domain: transmembrane #status predicted <TM4>
F:221-238/Domain: transmembrane #status predicted <TM5>
F:244-261/Domain: transmembrane #status predicted <TM6>
F:408-428/Domain: transmembrane #status predicted <TM7>

Query Match 61.9%; Score 1447; DB 2; Length 467;
Best Local Similarity 64.4%; Pred. No. 1.4e-100;
Matches 300; Conservative 41; Mismatches 83; Indels 42; Gaps 7;
Qy 24 ESPTPRSCQGRQPEDGENTAWRSQNEEDGEDDPRYVCSGVV-----GRPPG----- 75
Db 3 ELPAPLSYFQNAQSEDNHLSNTVRSQNDREQDGHDRRL-GNPELSNGRPOGNSGP 61
Qy 75 -----LEELTLKYGAHVIMLFVPTLCMVVVVATIKSVFRTEKNGQLIYTPFTE 127
Db 62 VVERDEEDELTLKYGAHVIMLFVPTLCMVVVVATIKSVFRTEKNGQLIYTPFTE 121
Qy 128 TPSVGORLLNSVNTLMISVIVMTIFLVLYKYRCYKFIHGWLMISLMFLFTYI 187
Db 122 TETVQGRALSHVLAAMISVIVMTIFLVLYKYRCYKFIHAWLIISLLEFFSYI 181
Qy 188 LGEVLKTYNVAMDYPTLLTVMNFGAVGVCIHMKGPLVLAQAYLIMISALMALYFIKYL 247
Db 182 LGEVFTYNAVADYITVALLIWNFGVGMISHWKGPLRLQAYLIMISALMALYFIKYL 241
Qy 248 PWSANWILGAISYVDLVAVLCPLKPLRMVETAGERNIPFALIYSAMVWVGMAKL 307
Db 242 PEWTAWLILAVISYVDLVAVLCPLKPLRMVETAGERNETLFPALIIYSTVWVLYNMAEG 301
Qy 308 DPSSQGLAL--QLPYD-----PEMEEDSYDSFGESYPVEFEP-----PLTGYPG 349
Db 302 DPEAQRVSKTKYNAQGTTEREAQASVPENDDGGFSEWEAQRDSQLGPHRSTSVSRAAV 361
Qy 350 EEL-----EEERGVKLGIDGFIYFSLVKGAAATGSGDWNNTLACFVAILGLCLT 402
Db 362 QEISSIPASEDPERGVKLGIDGFIYFSLVKGASATASGDWNTTIACFVAILGLCLT 421
Qy 403 LLLAVFKKALPALPISITFGILFYFTDNLVRPMDTLASHQLYI 448
Db 422 LLLAIFKKALPALPISITFGILFYFATDYLVPMDQLAFHOFYI 467

RESULT 10
JC5081
presenilin 1 protein isoform 463 - lesser mouse lemur
C:Species: Microcebus murinus (lesser mouse lemur)
C:Date: 31-Jan-1997 #sequence_revision 31-Jan-1997 #text_change 13-Sep-1998
C:Accession: JC5081
R:Calenda, A.; Mestre-Frances, N.; Czech, C.; Pradier, L.; Petter, A.; Bons, N.; Bellis,
presenilin 1, splice form 374 - human
N:Alternate names: Alzheimer's disease protein 3
C:Species: Homo sapiens (man)
C:Date: 20-Jul-1996 #sequence_revision 13-Mar-1997 #text_change 29-Sep-1999
C:Accession: S63684
R:Sahara, N.; Yahagi, Y.; Takagi, H.; Kondo, T.; Okochi, M.; Usami, M.; Shirasawa, T.
FEBS Lett. 381, 7-11, 1996
A:Title: Identification and characterization of presenilin I-467, I-463 and I-374.
A:Reference number: S63683; MUID:96193901
A:Accession: S63684

Biochem. Biophys. Res. Commun. 228, 430-439, 1996
A:Title: Molecular cloning, sequencing, and brain expression of the presenilin 1 gene
A:Reference number: JC5080; MUID:97079199
A:Contents: brain
A:Accession: JC5081
A:Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-463 <CAL>
A:Cross-references: EMBL:Z71333
C:Comment: This protein is an intermembrane protein with seven transmembrane domains.
C:Genetics:
A:Gene: psi
A:Map position: 14
C:Superfamily: presenilin
C:Keywords: transmembrane protein
F:78-96/Domain: transmembrane #status predicted <TM1>
F:129-150/Domain: transmembrane #status predicted <TM2>
F:160-181/Domain: transmembrane #status predicted <TM3>
F:191-209/Domain: transmembrane #status predicted <TM4>
F:217-234/Domain: transmembrane #status predicted <TM5>
F:240-257/Domain: transmembrane #status predicted <TM6>
F:404-424/Domain: transmembrane #status predicted <TM7>

Query Match 61.6%; Score 1440; DB 2; Length 463;
Best Local Similarity 64.2%; Pred. No. 4.7e-100;
Matches 300; Conservative 39; Mismatches 80; Indels 48; Gaps 8;
Qy 24 ESPTPRSCQGRQPEDG--ENTAWRSQNEED-----GEEDPDYVCSGVGPPG--- 75
Db 3 ELPAPLSYFQNAQSEDNHLSNTNREQDGHDRRLGNPELS-----NGRPOGNSG 56
Qy 75 -----LEELTLKYGAHVIMLFVPTLCMVVVVATIKSVFRTEKNGQLIYTPFTE 126
Db 57 PVVERDEEDELTLKYGAHVIMLFVPTLCMVVVVATIKSVFRTEKNGQLIYTPFTE 116
Qy 127 DPSPVGORLLNSVNTLMISVIVMTIFLVLYKYRCYKFIHGWLMISLMFLFTYI 186
Db 117 DFTVQGRALSHVLAAMISVIVMTIFLVLYKYRCYKFIHAWLIISLLEFFSFY 176
Qy 187 YLGEVLKTYNVAMDYPTLLTVMNFGAVGVCIHMKGPLVLAQAYLIMISALMALYFIKY 246
Db 177 YLGEVFTYNAVADYITVALLIWNFGVGMISHWKGPLRLQAYLIMISALMALYFIKY 236
Qy 247 LPWSANWILGAISYVDLVAVLCPLKPLRMVETAGERNIPFALIYSAMVWVGMAK 306
Db 237 LPWTAWLILAVISYVDLVAVLCPLKPLRMVETAGERNETLFPALIIYSTVWVLYNMAE 296
Qy 307 LDPSSQGLAL--QLPYD-----PEMEEDSYDSFGESYPVEFEP-----PLTGYP 348
Db 297 DPEAQRVSKTKYNAQGTTEREAQASVPENDDGGFSEWEAQRDSQLGPHRSTSVSRAA 356
Qy 349 GEEL-----EEERGVKLGIDGFIYFSLVKGAAATGSGDWNNTLACFVAILGLCL 401
Db 357 VQEISSIPASEDPERGVKLGIDGFIYFSLVKGASATASGDWNTTIACFVAILGLCL 416
Qy 402 TLLAVFKKALPALPISITFGILFYFTDNLVRPMDTLASHQLYI 448
Db 417 TLLAIFKKALPALPISITFGILFYFATDYLVPMDQLAFHOFYI 463

RESULT 11
S63684
presenilin 1, splice form 374 - human
N:Alternate names: Alzheimer's disease protein 3
C:Species: Homo sapiens (man)
C:Date: 20-Jul-1996 #sequence_revision 13-Mar-1997 #text_change 29-Sep-1999
C:Accession: S63684
R:Sahara, N.; Yahagi, Y.; Takagi, H.; Kondo, T.; Okochi, M.; Usami, M.; Shirasawa, T.
FEBS Lett. 381, 7-11, 1996
A:Title: Identification and characterization of presenilin I-467, I-463 and I-374.
A:Reference number: S63683; MUID:96193901
A:Accession: S63684

||||: : || | | : |
Db 336 YFSSHIALPFTDLCSQILILI 358

RESULT 14

T00724

presenilin homolog F22013.19 - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C:Date: 12-Feb-1999 #sequence_revision 12-Feb-1999 #text_change 20-Sep-1999
C:Accession: T00724
R:Shinn, P.; Buehler, E.; Dewar, K.; Peng, J.; Kim, C.; Li, Y.; Sun, H.; Conway, A.; Con-
eologis, A.; Ecker, J.R.
submitted to the EMBL Data Library, April 1998
A:Description: Genomic sequence for Arabidopsis thaliana BAC F22013.
A:Reference number: Z14200
A:Accession: T00724
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-453 <SHI>
A:Cross-references: EMBL:AC003981; NID:g3063438; PID:g3063457; GSPDB:GN0059; ATSP:F22013
C:Genetics:
A:Gene: ATSP:F22013.19
A:Map position: 1
A:Introns: 108/1
C:Superfamily: presenilin

Query Match 22.08; Score 513; DB 2; Length 453;

Best Local Similarity 31.1%; Pred. No. 5.2e-31;

Matches 136; Conservative 78; Mismatches 130; Indels 94; Gaps 12;

QY 88 VIMLFVPTLCMIVVATIKSVREYTEKN----GQLIYPTFTDTPSVGQRLLSVNLTL 143
Db 13 IIGVMAVPSICMELVLVLTSLSVTSDPQIRSAANLIYIENPSDSTTV--KLEGLSANAI 70
QY 144 IMISVIVMTIFVLVLYKYCYKFIHGLIMSLMLLFLEFYIYVLGEVLKTYNVAMDYPT 203
Db 71 VFVVLIAAVTIFVLVLYYNTFNELKHYMRSAFFVLGTMGGAIFLSIIQHFSIPVDSIT 130
QY 204 LLTVNFWGAVGMVCIHWK--PLVLOQAYLIMISALMALVFIKYLPEWSAWILGAISVY 262
Db 131 CFILLNFITLIGLSVFAGGPIVLROCYMVGMIVVAAWFK-LPEWTTWFIIVALALY 189
QY 263 DLVAVLCPKPLMLVETAQERNEPIPALIY-----SSAMVTVGMAKL 307
Db 190 DLVAVLAPGGLKLLVYELASSRDEEL-PAMVYEARTVSSGNQRNRGSSRLALVGGGV 248
QY 308 DPSOGALQLPYDEM----EEDSY-----DSFGPSYPVEFPEPLTGY 347
Db 249 SDGSVELQAVRHDVHVNQLGRENHNDYNAIVRDINDVDGIGNGSRGGLERSPLVGS 308
QY 348 PG-----EE-----LEEE-----ERGV 360
Db 309 PSASEHSTVGTGRNMDRESVDEEMSPVLGMWGDNRREARGLESNDVVDISNRGI 368
QY 361 KLGLGDFIFYSLVKGAAATSGDWNITLACFVAILIGLCTLLLLAVFKKALPALPISI 420
Db 369 KLGLGDFIFYSLVKGAAATSGDWNITLACFVAILIGLCTLLLLAVFKKALPALPISI 420
QY 421 TFLGLIFYSTDLNLRPFM 438
Db 426 MGVVFFLRLMEPFV 443

RESULT 15

A43459

sperm membrane protein spe-4 - Caenorhabditis elegans
N:Alternate names: probable integral membrane protein
C:Species: Caenorhabditis elegans
C:Date: 10-Jun-1993 #sequence_revision 18-Nov-1994 #text_change 09-Sep-1997
C:Accession: A43459; S24632; S24633
R:L'Hernault, S.W.; Arduengo, P.M.
J. Cell Biol. 119, 55-68, 1992

A:Title: Mutation of a putative sperm membrane protein in Caenorhabditis elegans prev
A:Reference number: A43459; MUID:92407040

A:Accession: A43459

A:Status: preliminary; not compared with conceptual translation

A:Molecule type: DNA; mRNA

A:Residues: 1-465 <LHE>

A:Cross-references: EMBL:Z14067; NID:g6868; PID:g6869; EMBL:Z14066; NID:g6870; PID:g6

A:Experimental source: strain Bristol N2

A:Note: the nucleotide sequence was submitted to the EMBL Data Library, July 1992

C:Genetics:

A:Introns: 69/3; 154/3; 200/1; 224/3; 300/1; 386/1; 435/1

Query Match 11.7%; Score 274; DB 2; Length 465;

Best Local Similarity 21.5%; Pred. No. 3.4e-13;

Matches 93; Conservative 80; Mismatches 127; Indels 132; Gaps 14;

QY 114 EKNGLIYPTFTEDT--PSVGORLINSVLN----LFTYIYVLGEVLKTYNVAMDYPTLL--TVNFGA 213
Db 42 EVNSELSTYFLDPSFEQTTGNLLLDGFIINGVGTILVIGCVSFIMLAF--VLEDFR--RI 97
QY 168 IHGWLIMSMLLF-----LFTYIYVLGEVLKTYNVAMDYPTLL--TVNFGA 213
Db 98 VKAWLTLSCLLILFGVSAQTLHDMFSQVFDQDDNQY-----YMTIVLIWVTVVYVGF- 152
QY 214 VGMVCIHWKGPLVLOQAYLIMISALMALVFIKYLPEWSAWILGAISVYDLVAVLCPKGP 273
Db 152 -GIYAFFSNSLLIHQIFVVTNCSLISVFLRVFPFSKTTFWLWILVFLWDLFAVLAPMGP 210
QY 274 LRMLVETAQERNEPIPALIYSSAMVTVGMAKLDPSSOG-----ALQLPYDPE 322
Db 211 LKKVQERASDYKCVLNLMFSANEKRLTAGSNQETNEGEESTIRRTVKQIETTYKRE 270
QY 323 MEEDSY-----DSF-----GEPSPYEPFPEPLTGYGPEEEEEEE--- 358
Db 271 AQDDEFYQKIRQRAAINPDSVPTEHSPIVEAEPSPIELKEKNST---EELSDDSDTS 326
QY 358 -----R 358
Db 327 ETSGSSSLSSSDSTTVSTSDISTAECDQKEDWDLVSNLSPNNDKRPATAADALNDGE 386
QY 359 GVKLGLGDFIFYSLVKGAAATSGDWNITLACFVAILIGLCTLLLLAVFKKALPALPI 418
Db 387 VRLUGFGDFIFYSLIGQAASGCP--FAVISAALGILFGLVVLTVFSTTESTTPALPL 444
QY 419 SITFGLIFYFST 430
Db 445 PVICGTCFCYFS 456

Search completed: March 18, 2000, 14:11:52

Job time: 489 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 16:08:11 ; Search time 32.57 Seconds
(without alignments)
410.791 Million cell updates/sec

Title: US-08-509-359B-137
Perfect score: 2336
Sequence: 1 MLTFMADSDSEEVCDERTSL.....STDNLVRFMDTLASHQLYI 448

Scoring table: BLOSUM62

Searched: 82229 seqs, 29864866 residues

Database : SwissProt_38:*

Word size : 0

Number of hits that pass the threshold : 82229

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	2336	100.0	448	1 PSN2_HUMAN	P49810 homo sapien
2	2232	95.5	448	1 PSN2_MOUSE	O61144 mus musculus
3	2227	95.3	448	1 PSN2_RAT	O88777 rattus norv
4	2218	94.9	445	1 PSN2_MICMU	P79801 microcebus
5	1624.5	69.5	449	1 PSN2_XENLA	O12977 xenopus lae
6	1468	62.8	467	1 PSN1_MOUSE	P49769 mus musculus
7	1467	62.8	467	1 PSN1_HUMAN	P49768 homo sapien
8	1459.5	62.5	468	1 PSN1_RAT	P97887 rattus norv
9	1449.5	62.1	433	1 PSN1_XENLA	O12976 xenopus lae
10	1447	61.9	467	1 PSN1_MICMU	P79802 microcebus
11	1150	49.2	541	1 PSN1_DROME	O02194 drosophila
12	1043	44.6	836	1 YLAK_CAEEL	Q20076 caenorhabdi
13	983	42.1	461	1 SE12_CAEEL	P52166 caenorhabdi
14	524.5	22.5	358	1 HOPI_CAEEL	O02100 caenorhabdi
15	513	22.0	453	1 PSNH_ARATH	O84668 arabidopsis
16	274	11.7	465	1 SPEX_CAEEL	Q01608 caenorhabdi
17	119	5.1	531	1 YDFG_SCHPO	Q10487 schizosacch
18	110	4.7	2016	1 CIN5_HUMAN	Q14524 homo sapien
19	109	4.7	1840	1 CIN4_RAT	P15390 rattus norv
20	108	4.6	381	1 CYB_DASCR	Q34302 dasyercus
21	104	4.5	1476	1 CFTR_MOUSE	P26361 mus musculus
22	103.5	4.4	1450	1 CFTR_RABIT	Q00554 oryctolagus
23	103	4.4	382	1 CYB_DIDMA	P41303 didelphis m
24	103	4.4	380	1 CYB_MICLO	P56731 microtus lo
25	102.5	4.4	893	1 BOSS_DROVI	Q24738 drosophila
26	102.5	4.4	381	1 CYB_ANTFL	Q33706 antechinus
27	102	4.4	381	1 CYB_DASMA	Q03522 dasyurus ma
28	102	4.4	381	1 CYB_PSENI	P35553 pseudantech
29	101.5	4.3	381	1 CYB_NINIV	Q35196 ningau yvo
30	101.5	4.3	381	1 CYB_PARAP	Q35377 parantechin
31	101	4.3	381	1 CYB_PLAMS	Q35533 planigale m
32	100.5	4.3	1951	1 CIN3_RAT	P08104 rattus norv
33	100.5	4.3	381	1 CYB_PSEMD	O03543 pseudantech
34	100	4.3	381	1 CYB_DASGE	O20604 dasyurus ge
35	99	4.2	2019	1 CIN5_RAT	P15389 rattus norv
36	99	4.2	381	1 CYB_ANTMI	O63534 antechinus
37	99	4.2	381	1 CYB_PHATA	Q35673 phascogale
38	98.5	4.2	872	1 CIQ3_HUMAN	O43525 homo sapien
39	98	4.2	1836	1 CIN4_HUMAN	P35499 homo sapien

40	98	4.2	381	1	CYB_ANTME	Q33782 antechinus
41	98	4.2	381	1	CYB_DASAL	Q34289 dasyurus al
42	98	4.2	381	1	CYB_SMICR	Q35810 smanthopsis
43	98	4.2	460	1	NU4M_GADMO	P55781 gadus morhu
44	98	4.2	217	1	YPRA_ECOLI	P13974 escherichia
45	97.5	4.2	381	1	CYB_DASHA	Q34321 dasyurus ha

ALIGNMENTS

RESULT 1				
PSN2_HUMAN				
ID	PSN2_HUMAN	STANDARD;	PRT;	448 AA.
AC	P49810;			
DT	01-OCT-1996	(Rel. 34, Created)		
DT	01-OCT-1996	(Rel. 34, Last sequence update)		
DT	15-JUL-1999	(Rel. 38, Last annotation update)		
DE	PRESENTLIN 2 (PS-2) (STM-2) (E5-1) (AD3LP) (AD5).			
GN	PSEN2 OR PSN2 OR AD4 OR PS2 OR STM2.			
OS	Homo sapiens (Human).			
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia;			
CC	Eutheria; Primates; Catarrhini; Hominidae; Homo.			
RN	[1]			
RP	SEQUENCE FROM N.A., AND VARIANT FAD ILE-141.			
RX	MEDLINE; 95365816.			
RA	LEVY-LAHAD E., WASSCO W., POORRAJ P., ROMANO D.M., OSHIMA J.,			
RA	PETTINGELL W.H., YU C.-E., JONDRO P.D., SCHMIDT S.D., WANG K.,			
RA	CROWLEY A.C., FU Y.-H., GUENETTE S.Y., GALAS D., NEMENS E.,			
RA	WIJSMAN E.M., BIRD T.D., SCHELLENBERG G.D., TANZI R.E.;			
RT	"Candidate gene for the chromosome 1 familial Alzheimer's disease			
RT	locus.";			
RL	Science 269:973-977(1995).			
RN	[2]			
RP	SEQUENCE FROM N.A., AND VARIANTS FAD ILE-141 AND VAL-239.			
RC	TISSUE-BRAIN, AND COLON;			
RX	MEDLINE; 95379971.			
RA	ROGAEV E.I., SHERRINGTON R., ROGAEVA E.A., LEVESQUE G., IKEDA M.,			
RA	LIANG Y., CHI H., LIN C., HOLMAN K., TSUDA T., MAR L., SORBI S.,			
RA	NACMIAS B., PIACENTINI S., AMADUCCI L., CHUMAKOV I., COHEN D.,			
RA	LANNFELT L., FRASER P.E., ROMMENS J.M., ST GEORGE-HYSLOP P.H.;			
RT	"Familial Alzheimer's disease in kindreds with missense mutations in			
RT	a gene on chromosome 1 related to the Alzheimer's disease type 3			
RT	gene.";			
RL	Nature 376:775-778(1995).			
RN	[3]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE; 96109229.			
RA	LI J., MA J., POTTER H.;			
RT	"Identification and expression analysis of a potential familial			
RT	Alzheimer disease gene on chromosome 1 related to AD3.";			
RL	Proc. Natl. Acad. Sci. U.S.A. 92:12180-12184(1995).			
RN	[4]			
RP	SEQUENCE FROM N.A.			
RA	LEVY-LAHAD E., POORRAJ P., WANG K., FU Y.H., OSHIMA J.,			
RA	MULLIGAN J., SCHELLENBERG G.D.;			
RL	Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.			
RN	[5]			
RP	REVIEW ON VARIANTS.			
RX	MEDLINE; 98180715.			
RA	CRUTS M., VAN BROECKHOVEN C.;			
RT	"Presenilin mutations in Alzheimer's disease.";			
RL	Hum. Mutat. 11:183-190(1998).			
RN	[6]			
RP	VARIANT AD HIS-62.			
RX	MEDLINE; 98046005.			
RA	CRUTS M., VAN DUJN C.M., BACKHOVENS H., VAN DEN BROECK M.,			
RA	WEHNET A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J.,			
RA	ST GEORGE-HYSLOP P.H., HOFMAN A., VAN BROECKHOVEN C.;			
RT	"Estimation of the genetic contribution of presenilin-1 and -2			
RT	mutations in a population-based study of presenile Alzheimer			
RT	disease.";			
RL	Hum. Mol. Genet. 7:43-51(1998).			

```
DR EMBL; U57325; AAC53311.1; -.
DR EMBL; AF038935; AAB92660.1; -.
DR EMBL; U49111; AAC52935.1; -.
DR MGD; MGI:109284; PSEN2.
DR PFAM; PF01080; Presenilin; 1.
KW Transmembrane; Alternative initiation.
FT CHAIN 1 448 PRESENILIN 2.
FT CHAIN 298 448 PRESENILIN 2-SHORT.
FT TRANSMEM 88 106 POTENTIAL.
FT TRANSMEM 142 160 POTENTIAL.
FT TRANSMEM 167 188 POTENTIAL.
FT TRANSMEM 203 219 POTENTIAL.
FT TRANSMEM 230 246 POTENTIAL.
FT TRANSMEM 253 269 POTENTIAL.
FT TRANSMEM 288 305 POTENTIAL.
FT TRANSMEM 387 406 POTENTIAL.
FT TRANSMEM 413 429 POTENTIAL.
FT CONFLICT 87 87 R -> H (IN REF. 2).
FT CONFLICT 226 226 A -> V (IN REF. 2).
FT CONFLICT 324 324 MISSING (IN REF. 2).
SQ SEQUENCE 448 AA; 49955 MW; 680ACF19 CRC32;

Query Match 95.5%; Score 2232; DB 1; Length 448;
Best Local Similarity 95.5%; Pred. No. 7 3e-147;
Matches 428; Conservative 6; Mismatches 14; Indels 0; Gaps 0;

QY 1 MLTFMSDSEEEVCDERTSLMSAESPTRSCQEGRQGPEDGENTAQRSENEDEEDP 60
DB 1 MLAFMSDSEEEVCDERTSLMSAESPTRSCQEGRQGPEDGENTAQRSENEDEEDP 60
QY 61 DRYCVCVGPGRPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLI 120
DB 61 DRYACSGAPGRPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLI 120
QY 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLMSSMLL 180
DB 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLMSSMLL 180
QY 181 FLFTYIYLGEVLKTYNVAMDYPTLLFVWVNFAGVMYCIHWKGLVPLVQAYLIVISALMA 240
DB 181 FLFTYIYLGEVLKTYNVAMDYPTLLFVWVNFAGVMYCIHWKGLVPLVQAYLIVISALMA 240
QY 241 LVPIKILPEWSAVILGAISYDVLVAVLCRPGPLMLVETAQERNEIFPALIYSSAMVW 300
DB 241 LVPIKILPEWSAVILGAISYDVLVAVLCRPGPLMLVETAQERNEIFPALIYSSAMVW 300
QY 301 TVGMAKLDPSQCALQPLYPMEEDSYDSFGPSYPEVEPPLTGYPGLEEEERGV 360
DB 301 TVGMAKLDPSQCALQPLYPMEEDSYDSFGPSYPEVEPPLTGYPGLEEEERGV 360
QY 361 KLGLGDFIFYSVLVGAAATGSGDWNNTLACFAVAILGLCITLLLLAVFKKALPALPISI 420
DB 361 KLGLGDFIFYSVLVGAAATGSGDWNNTLACFAVAILGLCITLLLLAVFKKALPALPISI 420
QY 421 TFGLIIFYSTDLNLRPMDTFLASHQLYI 448
DB 421 TFGLIIFYSTDLNLRPMDTFLASHQLYI 448

RESULT 3
PSN2_RAT
ID PSN2_RAT STANDARD; PRT; 448 AA.
AC O88777; O35546; O08947;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN 2 (PS-2).
GN Rattus norvegicus (Rat).
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
RN [1]

SEQUENCE FROM N.A.
STRAIN-WISTAR; TISSUE-BRAIN;
PRENTZEL S., ABDEL A.S., LUEBBERT H.;
Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
[2]
SEQUENCE FROM N.A.
STRAIN-WISTAR; TISSUE-BRAIN;
MEDLINE; 97473536.
TAKAHASHI H., MERCKEN M., NAKAZATO Y., NOGUCHI K., MURAYAMA M.,
IMAHORI K., TAKASHIMA A.;
"Cloning of cDNA and expression of the gene encoding rat
presenilin-2.";
Gene 197:383-387(1997).
[3]
SEQUENCE FROM N.A.
STRAIN-WISTAR; TISSUE-BRAIN;
MEDLINE; 98207716.
TANAHASHI H., TABIRA T.;
"Cloning of the cDNA encoding rat presenilin-2.";
Biochim. Biophys. Acta 1396:259-262(1998).
RT FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE
EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE. MAY
FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS (BY
SIMILARITY).
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
CC
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-----
EMBL; X93267; CAA67663.1; -.
EMBL; D83700; BAA2832.1; -.
EMBL; AB004454; BAA20406.1; -.
DR PFAM; PF01080; Presenilin; 1.
DR TRANSMEM 88 106 POTENTIAL.
DR TRANSMEM 142 160 POTENTIAL.
DR TRANSMEM 167 188 POTENTIAL.
DR TRANSMEM 203 219 POTENTIAL.
DR TRANSMEM 230 246 POTENTIAL.
DR TRANSMEM 253 269 POTENTIAL.
DR TRANSMEM 288 305 POTENTIAL.
DR TRANSMEM 387 406 POTENTIAL.
DR TRANSMEM 413 429 POTENTIAL.
DR CONFLICT 7 7 S -> T (IN REF. 1).
DR CONFLICT 86 87 KH -> ND (IN REF. 3).
SQ SEQUENCE 448 AA; 50051 MW; E80DF681 CRC32;

Query Match 95.3%; Score 2227; DB 1; Length 448;
Best Local Similarity 95.3%; Pred. No. 1.6e-146;
Matches 427; Conservative 6; Mismatches 15; Indels 0; Gaps 0;

QY 1 MLTFMSDSEEEVCDERTSLMSAESPTRSCQEGRQGPEDGENTAQRSENEDEEDP 60
DB 1 MLTFMSDSEEEVCDERTSLMSAESPTRSCQEGRQGPEDGENTAQRSENEDEEDP 60
QY 61 DRYCVCVGPGRPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLI 120
DB 61 DRYACSGVGPGRPGLEELTKYGAHVIMLFVPTLCMIVVATIKSVRYTEKNGQLI 120
QY 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLMSSMLL 180
DB 121 YTPFTEDTPSVGQRLNSVLTLMISIVVMTIFLVLYKYRCYKFIHGWLMSSMLL 180
QY 181 FLFTYIYLGEVLKTYNVAMDYPTLLFVWVNFAGVMYCIHWKGLVPLVQAYLIVISALMA 240
DB 181 FLFTYIYLGEVLKTYNVAMDYPTLLFVWVNFAGVMYCIHWKGLVPLVQAYLIVISALMA 240
```


RA ROYTA M., LILIUS L., EEROLA A., ST GEORGE-HYSLOP P.H., FREY H.,
 RA LANNFELT L.,
 RT "The Glu318gly mutation of the presenilin-1 gene does not necessarily
 cause Alzheimer's disease";
 RL Ann. Neurol. 44:965-967(1998).
 RN [15]
 RP VARIANT GLY-318.
 RX MEDLINE; 99066775.
 RA ALDUDO J., BULLIDO M.J., FRANK A., VALDIVIESO F.;
 RT "Missense mutation E318G of the presenilin-1 gene appears to be a
 nonpathogenic polymorphism";
 RL Ann. Neurol. 44:985-986(1998).
 RN [16]
 RP VARIANTS AD VAL-79; CYS-115; VAL-231, AND VARIANT GLY-318.
 RX MEDLINE; 98046005.
 RA CRUTS M., VAN DOIJN C.M., BROCKHOVEN H., VAN DEN BROECK M.,
 RA WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J.,
 RA ST GEORGE-HYSLOP P.H., HOFMAN A., VAN BROECKHOVEN C.;
 RT "Estimation of the genetic contribution of presenilin-1 and -2
 mutations in a population-based study of presenile Alzheimer
 disease";
 RL Hum. Mol. Genet. 7:43-51(1998).
 RN [17]
 RP VARIANTS AD D-120; R-163; V-209; V-260; L-264; Y-410 AND P-426.
 RX MEDLINE; 98180720.
 RA POORAJ P., SHARMA V., ANDERSON L., NEMENS E., ALONSO M.E., ORR H.,
 RA WHITE J., HESTON L., BIRD T.D., SCHELEBERG G.D.;
 RT "Missense mutations in the chromosome 14 familial Alzheimer's disease
 presenilin 1 gene";
 RL Hum. Mutat. 11:216-221(1998).
 RN [18]
 RP VARIANT AD GLU-378.
 RX MEDLINE; 99211215.
 RA BESANCON R., LORENZI A., CRUTS M., RADAWIEC S., STURTZ F.,
 RA BROUSSOLLE E., CHAZOT G., VAN BROECKHOVEN C., CHAMBA G.,
 RA VANDENBERGHE A.;
 RT "Missense mutation in exon 11 (codon 378) of the presenilin-1 gene in
 a French family with early-onset Alzheimer's disease and transmission
 study by mismatch enhanced allele specific amplification";
 RL Hum. Mutat. 11:481-481(1998).
 RN [19]
 RP VARIANTS AD LEU-169 AND GLN-436.
 RX MEDLINE; 99047368.
 RA TADDEI K., KWOK J.B., KRIL J.J., HALLIDAY G.M., CREASEY H.,
 RA HALLUPP M., FISHER C., BROOKS W.S., CHUNG C., ANDREWS C.,
 RA MASTERS C.L., SCHOFIELD P.R., MARTINS R.N.;
 RT "Two novel presenilin-1 mutations (Ser169Leu and Pro436Gln) associated
 with very early onset Alzheimer's disease";
 RL NeuroReport 9:3335-3339(1998).
 RN [20]
 RP VARIANT AD PRO-169.
 RX MEDLINE; 99148656.
 RA EZQUEERRA M., CARNERO C., BLESER R., GELPI J.L., BALLESTA F., OLIVA R.;
 RT "A presenilin 1 mutation (Ser169Pro) associated with early-onset AD
 and myoclonic seizures";
 RL Neurology 52:566-570(1999).

Query Match 62.8%; Score 1467; DB 1; Length 467;
 Best Local Similarity 65.7%; Pred. No. 3.5e-94;
 Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;

QY 24 ESTPTSCQGRGPDGENTAWRSQNEDEEDDPRVCVSGVP-----GRPPG---- 75
 DB 3 ELPAFLSYFQNAQMSDENHLSNTVRSQNDNRQEHNDNR-SLGHPEPLNGRPOGNSRQ 61
 QY 75 -----LEELTKYGAKIVMLFVPTLCMVVATIKSVFYTEKNGQLIYPTPTD 127
 DB 62 VFGQDEEELTKYGAKIVMLFVPTLCMVVATIKSVFYTEKNGQLIYPTPTD 121
 QY 128 TPVSGQRLLNSLTMISVIVVMTIFLVLYKRYCKFIHGWLIMSSMLLFLFTYIY 187
 DB 122 TEIVGQRALHSILNAATMISVIVVMTILLVLYKRYCKVIHAWLISSLLLLFFFSFIY 181

QY 188 LGEVLKTYNTVAMDYPTLLLTWNFGAVGVMCIHWKGLVPLQQAYLIMISALMALVFIKYL 247
 DB 182 LGEVFKTYNTVADYITVALLIWNFGVGMISHWKGLRQOAYLIMISALMALVFIKYL 241
 QY 248 PEWSAWVILGAIISYDVLVAVLCPRKPLMLVETAQERNEPIFPALIISSAMVTVGMAKL 307
 DB 242 PEWTAWLILAVISYDVLVAVLCPRKPLMLVETAQERNETLFPALIISSAMVTVGMVNAEG 301
 QY 308 DPSSQGLAL--QLPYDPE-MEEDSDYDSFGE---PSYPEVFEPPLTGYPG----- 350
 DB 302 DPEAQREVSKNSKYNASTERESQDTVAENDDGGFSEWEAQQRDSHLGPHRSTPESRAAV 361
 QY 350 EEL-----EEEEERGKVLGDFIFYSVLVGAATGSGDWNNTTACFAVAILIGLCLT 402
 DB 362 QELSSSILAGEDPEERGKVLGDFIFYSVLVGRASATASGDWNNTTACFAVAILIGLCLT 421
 QY 403 LLLAVFKKALPALPISITITGLFIYFSTDLNLRPFMDTLASHQLYI 448
 DB 422 LLLAIFKKALPALPISITITGLFIYFATDYLQVFPMDQLAFHQFYI 467
 RESULT 8
 PSNL_RAT
 ID PSNL_RAT STANDARD; PRT; 468 AA.
 AC P97887; P97529;
 DT 15-JUL-1999 (Rel. 38, Created)
 DT 15-JUL-1999 (Rel. 38, Last sequence update)
 DE PRESENILIN 1 (PS-1) (S182 PROTEIN).
 GN PSEN1 OR PSNL1.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-WISTAR; TISSUE-BRAIN;
 RX MEDLINE; 97199371.
 RA TANIGUCHI T., HASHIMOTO T., TANIGUCHI R., SHIMADA K., KAWAMATA T.,
 RA YASUDA M., NAKAI M., TERASHIMA A., KOIZUMI T., MAEDA K., TANAKA C.;
 RT "Cloning of the cDNA encoding rat presenilin-1";
 RL Gene 186:73-75(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN-WISTAR; TISSUE-BRAIN;
 RX MEDLINE; 9625262.
 RA TAKAHASHI H., MURAYAMA M., TAKASHIMA A., MERCKEN M., NAKAZATO Y.,
 RA NOGUCHI K., IMAHORI K.;
 RT "Molecular cloning and expression of the rat homologue of
 presenilin-1";
 RL Neurosci. Lett. 206:113-116(1996).
 CC -!- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE
 EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE.
 CC MAY FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS.
 CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
 CC -!- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
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 or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; D82578; BAA11575.1; -;
 DR EMBL; D82363; BAA11564.1; -;
 DR PFAM; PF01080; Presenilin; 1.
 KW Transmembrane; Glycoprotein.
 FT TRANSMEM 83 103 POTENTIAL.
 FT TRANSMEM 133 153 POTENTIAL.
 FT TRANSMEM 161 181 POTENTIAL.
 FT TRANSMEM 195 215 POTENTIAL.
 FT TRANSMEM 221 241 POTENTIAL.

PSN1_MICMU STANDARD; PRT; 467 AA.
AC P79802;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN 1 (PS-1).
GN PSN1 OR PSN1 OR PS1.
OS Microcebus murinus (Lesser mouse lemur).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Strepsirhini; Cheirogaleidae; Microcebus.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-BRAIN;
RX MEDLINE: 97079199.
RA CALEDA A., MESTRE-FRANCES N., CZECH C., PRADIER L., BONS N.,
RA BELLIS M.;
RT "Molecular cloning, sequencing, and brain expression of the
RT presenilin 1 gene in Microcebus murinus.";
RL Biochem. Biophys. Res. Commun. 228:430-439(1996).
CC -1- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE
CC EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE.
CC MAY FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -1- ALTERNATIVE PRODUCTS: TWO ISOFORMS: I-467 (SHOWN HERE) AND I-
CC 463; ARE PRODUCED BY ALTERNATIVE SPLICING.
CC -1- TISSUE SPECIFICITY: FOUND PREDOMINANTLY IN NEURONS OF THE
CC DIFFERENT CORTICAL LAYERS AND HIPPOCAMPUS BUT ALSO IN SUBCORTICAL
CC STRUCTURES.
CC -1- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: Z71333; CA95930.1; -;
DR PFAM: PF01080; Presenilin; 1.
KW Transmembrane; Alternative splicing; Glycoprotein.
FT TRANSMEM 83 103 POTENTIAL.
FT TRANSMEM 133 153 POTENTIAL.
FT TRANSMEM 161 181 POTENTIAL.
FT TRANSMEM 191 211 POTENTIAL.
FT TRANSMEM 221 241 POTENTIAL.
FT TRANSMEM 244 264 POTENTIAL.
FT TRANSMEM 281 301 POTENTIAL.
FT TRANSMEM 408 428 POTENTIAL.
FT TRANSMEM 433 453 POTENTIAL.
FT CARBOHYD 279 279 POTENTIAL.
FT CARBOHYD 405 405 POTENTIAL.
FT VARSPLIC 26 29 MISSING (IN ISOFORM I-463).
SQ SEQUENCE 467 AA; 52384 MW; A841A0B7 CRC32;

Query Match 61.9%; Score 1447; DB 1; Length 467;
Best Local Similarity 64.4%; Pred. No. 8.4e-93;
Matches 300; Conservative 41; Mismatches 83; Indels 42; Gaps 7;

QY 24 ESPTPRSCGRCQGPGEQNTAQRWSENEDGEDDDRYVCSGVP-----GRPPG----- 75
Db 3 ELPAFLSYFQAQMSQNDHLSNTVRQNDREQQDGHDRRL-GNPEPLSNGRPQNGSGP 61

QY 75 -----LEELTKYGAHVIMLFPVPTLCMVIVVATIKSVRETKNGOLIYTPPTED 127
Db 62 VVERDEEDELTKYGAHVIMLFPVPTLCMVIVVATIKSVRYTRKDGOLIYTPPTED 121

QY 128 TPSVQRLNLSLTLTMSIVVMTIFLVLYKYRCYKFTGHWLMSLSLMLFLFTYIY 187
Db 122 TEITVQGRALHSVLNAATMSIVVMTILLVLYKYRCYKVIHAWLISSLLLLFFFSFIY 181

QY 188 LGEVLKTYNVAMDYPTLLLTWNFGAVGMVCIHWKGPLVLQOAYLIMISALMALVFIKYL 247
Db 182 LGEVFKTYNVAVDYITVALLTWNFGVGMISIHKGPLRLQOAYLIMISALMALVFIKYL 241
QY 248 PWSAWILGAIISYDLVAVLCPKGPLRLMVLVETAOERNEPIFPALIIYSAMVYTMGAKL 307
Db 242 PEWTAWLILAVISYDLVAVLCPKGPLRLMVLVETAOERNEPIFPALIIYSSTMYLVNMAEG 301
QY 308 DPSQOGAL--OLPYD-----PEMEEDSDVSFGEPSYPEVPEP----PLTGYPS 349
Db 302 DPEAQRVSKNTKYNAOQTEREAQASVPENDGGFSEWEAQORDSQLGPHRSTSVSRAV 361
QY 350 EEL-----EEERGVKLGDFIFYSVLVGRKAAATGSDWNTTLACFVAILGLCLT 402
Db 362 QEISSSPASEDPEERGKVLGDFVYFVSVLVGRASATASGDWNTTLACFVAILGLCLT 421
QY 403 LLLLAFFKALPALPISITIFGLIYFSTDLNVRPFMDTIALSHQLYI 448
Db 422 LLLLAFFKALPALPISITIFGLVFFATDYLQVPMQDLAFHOFYI 467

RESULT 11
PSN1_DROME STANDARD; PRT; 541 AA.
AC O02194; O02395; O76802;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE PRESENILIN HOMOLOG (DPS) (DMPS).
GN PS.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
RN [1]
RX SEQUENCE FROM N.A. (LONG ISOFORM).
RX MEDLINE: 97285868.
RA BOULIANNE G.L., LYNE-BAR I., HUMPHREYS J.M., LIANG Y., LIN C.,
RA ROGAEV E., ST GEORGE-HYSLOP P.;
RT "Cloning and characterization of the Drosophila presenilin
RT homologue.";
RL NeuroReport 8:1025-1029(1997).
RN [2]
RP SEQUENCE FROM N.A. (SHORT ISOFORM).
RC STRAIN-CANTON-S;
RA HONG C.S., KOO E.H.;
RT "Isolation and characterization of Drosophila presenilin homolog.";
RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A. (LONG AND SHORT ISOFORMS).
RA YE Y., FORTINI M.E.;
RT "Characterization of Drosophila presenilin and its colocalization
RT with Notch during development.";
RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
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CC -----
DR EMBL: U77934; AAB61139.1; -;
DR EMBL: U78084; AAB53369.1; -;
DR EMBL: AF084184; AAC33129.1; -;
DR EMBL: AF084184; AAC33128.1; -;
DR FLYBASE: FBgn0019947; PS.
DR PFAM: PF01080; Presenilin; 1.
KW Transmembrane; Glycoprotein; Alternative splicing.
FT TRANSMEM 107 127

```

Query Match      42.1%; Score 983; DB 1; Length 461;
Best Local Similarity 49.1%; Pred. No. 7.3e-61;
Matches 222; Conservative 60; Mismatches 92; Indels 78; Gaps 12;

QY 26 PTPRSQOEGRQGPEDGE-NTAQWRSQSNEDEGDDPRYVCSGVGPRPPGLEELTKYG 84
   : : | | | | | : : : | | : | | |
Db 2 PSTRRQOEG--GGADAETHVYGTNLITNRNSQDEN-----VVAEALKYG 46
   : | | | | | : : : | | : | | |

QY 85 AKHVMILFVPVTCMIVVWVATIKSVRYETKNGQ-LIYTPFTEPTPSVGORLLNSVLNTL 143
   | | | | | | | | | : : : | | : | | | : | : | : | : |
Db 47 ASHVILHUFVPSLCMALVUFTMTNITITSQNGRHLLYTPFVRETDTSIVK6KLSLGNAL 106
   : | | | | | | | | | : : : | | : | | : | : | : | : |

QY 144 IMISVIVMVTIFLVLYKRYCYAFIHGWLIMSSLMMLLFETYIYVLGVLTNTYNVAMDYPT 203
   : : | : | : | : | | | | | : | | | | | : | : | : | : |
Db 107 VMLCVVVLMTVLLIVFYKRYFYKLHGWLIVSVSFLLLFLETTYVQVFLSFDVSPSALL 166
   : | : | | | | | | | | | | | | | | | | | | | | : | : | : |

QY 204 LLITVWNGAVGWCVHWKGPLVLQQAYLIMISALMALVFIKYLPWSAWVLGAISYVD 263
   : | : | : | | | | | | | | | | | | | | | | | | : | : | : |
Db 167 VLFGLNGYVGLGMMCIHWKGPLRQOQFYLTMTSALMALVFIKYLPWTVMVFLVFSVWD 226
   : | : | | | | | | | | | | | | | | | | | | | | : | : | : |

QY 264 LVAVLCPKGPLMLIVETAQRNEPIFFALYISSMWVTVGMAKLDPSQOALQLPYDEM 323
   | | | | | | | | | | | | | | | | | : : : | : | : | : |
Db 227 LVAVLTPKGLRLVETAQRNEPIFFALYISSGVITYPYLVLT-----AVENTTDP- 279
   | | | | | | | | | | | | | | | | | : : : | : | : | : |

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CC      CC      EMBL; AF021905; AAB84394.1; -.
CC      CC      EMBL; AF000265; AAB52948.1; -.
CC      CC      WORMPEP; C18E3.8; CE08317.
CC      CC      PRAM; PF01080; Presenilin; 1.
CC      CC      Transmembrane.
CC      CC      KW      Transmem 13      33      POTENTIAL.
CC      CC      FT      Transmem 58      78      POTENTIAL.
CC      CC      FT      Transmem 87      107     POTENTIAL.
CC      CC      FT      Transmem 116     136     POTENTIAL.
CC      CC      FT      Transmem 149     169     POTENTIAL.
CC      CC      FT      Transmem 171     191     POTENTIAL.
CC      CC      FT      Transmem 298     318     POTENTIAL.
CC      CC      FT      Transmem 322     342     POTENTIAL.
CC      CC      SQ      SEQUENCE 358 AA; 39864 MW; FC6305ED CRC32;
CC      CC      Query Match 22.5%; Score 524.5; DB 1; Length 358;
CC      CC      Best Local Similarity 30.5%; Pred. No. 2.1e-29;
CC      CC      Matches 117; Conservative 81; Mismatches 136; Indels 49; Gaps 8;
CC      CC      QY      83 YGAKHVIMLFPVTLICMIVVATIKSVRFYTEKNGQLIYTPF-----TEDTPSVGQRLLNS 138

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GenCore version 4.5
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OM protein - protein search, using sw model
Run on: March 18, 2000, 22:05:40 ; Search time 49.26 seconds
(without alignments)
630.566 Million cell updates/sec

Title: US-08-509-359B-137
Perfect score: 2336
Sequence: 1 MLTFMADSEEEVCDERTSL.....STDNLVRPFMDTLASHQLYI 448

Scoring table: BLOSUM62

Searched: 225878 seqs, 69334122 residues

Database : SPTREMBL_12.*

Word size : 0

Number of hits that pass the threshold : 225878

- 1: sp_archaea.*
- 2: sp_bacteria.*
- 3: sp_fungi.*
- 4: sp_human.*
- 5: sp_invertebrate.*
- 6: sp_mammal.*
- 7: sp_mhc.*
- 8: sp_organelle.*
- 9: sp_phage.*
- 10: sp_plant.*
- 11: sp_rodent.*
- 12: sp_virus.*
- 13: sp_vertebrate.*
- 14: sp_unclassified.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	2262.5	96.9	449	6 Q9XT96	Q9xt96 bos taurus
2	1438.5	61.6	478	6 Q9XT97	Q9xt97 bos taurus
3	1407.5	60.3	456	13 Q9W6T7	Q9w6t7 brachydanio
4	1355	58.0	384	13 Q73869	Q73869 cyprinus ca
5	576.5	24.7	272	5 Q96340	Q96340 drosophila
6	440	18.8	184	4 Q95465	Q95465 homo sapien
7	113.5	4.9	406	5 Q19737	Q19737 caenorhabdi
8	112.5	4.8	4578	13 Q42181	Q42181 fugu rubrip
9	110.5	4.7	320	8 Q34086	Q34086 coccyzus er
10	110.5	4.7	381	8 Q35425	Q35425 phascoloror
11	109	4.7	1840	11 Q70611	Q70611 rattus norv
12	108.5	4.6	380	8 Q92209	Q92209 upupa epops
13	107.5	4.6	748	2 Q32577	Q32577 streptomyce
14	107	4.6	381	8 Q33723	Q33723 antechinus
15	106.5	4.6	382	8 Q34340	Q34340 didelphis m
16	106.5	4.6	318	11 Q35294	Q35294 rattus norv
17	106.5	4.6	303	11 P97829	P97829 rattus norv
18	106	4.5	652	5 Q93346	Q93346 caenorhabdi
19	106	4.5	382	8 Q34279	Q34279 didelphis a
20	106	4.5	382	8 Q34677	Q34677 gliroia ve
21	106	4.5	1717	13 Q90519	Q90519 fugu rubrip
22	105	4.5	444	2 Q9X2N3	Q9x2n3 arthrobacte
23	105	4.5	379	8 Q34428	Q34428 echimys did
24	104	4.5	382	8 Q35561	Q35561 phillander o
25	104	4.5	379	8 Q36096	Q36096 trinomys pa

ALIGNMENTS

RESULT 1

Q9XT96 PRELIMINARY; PRT; 449 AA.
AC Q9XT96;
DT 01-NOV-1999 (Tremblrel. 12, Created)
DT 01-NOV-1999 (Tremblrel. 12, Last sequence update)
DT 01-NOV-1999 (Tremblrel. 12, Last annotation update)
DE PRESENILIN 2.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovinae; Bos.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=BRAIN;
RA SAHARA N., SHIRASAWA T., MORI H.;
RT "Molecular cloning of bovine presenilin 2 gene";
RL Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF038937; AAD39024.1; -;
SQ SEQUENCE 449 AA; 50301 MW; A3DA878F CRC32;

Query Match	96.9%	Score 2262.5;	DB 6;	Length 449;
Best Local Similarity	97.1%	Pred. No. 4.7e-162;		
Matches 436;	Conservative 5;	Mismatches 7;	Indels 1;	Gaps 1;
QY 1	MLTFMADSEEEVCDERTSLMSAESPTPRSCQGRGPDGENTAWRSQENEED-GEED	59		
Db 1	MLTFMADSEEEVCDERTSLMSAESPTPRSCQGRGLEDGESAAQWRSQSEDEHEED	60		
QY 60	PRYVCSGVGPGRPPGLEELTLKYAKHVMFLFVPTVLCMVVVVATIKSVRFYTEKNGQL	119		
Db 61	PRYVCSGVGPGRPPGLEELTLKYAKHVMFLFVPTVLCMVVVVATIKSVRFYTEKNGQL	120		
QY 120	ITYPTFEDPSPGQRLNLSVNTLIMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLM	179		
Db 121	ITYPTFEDPSPGQRLNLSVNTLIMISIVVMTIFLVLYKYRCYKFIHGWLIMSSLM	180		
QY 180	LELFYIYLGEVLKYNVAMDPTLLTVNFGAVGMVCIHWKGPLVLOQAYLIMISALM	239		
Db 181	LELFYIYLGEVLKYNVAMDPTLLTVNFGAVGMVCIHWKGPLVLOQAYLIMISALM	240		
QY 240	ALVFIKYLPEWSAWTILGAISVYDLVAVLCPKGPLRMLVETAQERNEPIFALIYSSAMV	299		
Db 241	ALVFIKYLPEWSAWTILGAISVYDLVAVLCPKGPLRMLVETAQERNEPIFALIYSSAMV	300		
QY 300	WTVGMAKLDPSQGAQLQLPDPEMEEDSDTSDFCEPSYPEVFEPPLTGYPGCEELEBERG	359		

FN SEQUENCE FROM N.A.
RA ARCHER S., HIRANO J., DISS J.K., FRASER S.P., DJANGOZ M.B.A.;
RL NeuroReport 0:0-0(0).
DR EMBL; Y17128; CAA76641.1; -.
DR PFAM; YF01080; Presenilin; 1.
FT NON_TER 1
SQ SEQUENCE 384 AA; 43276 MW; 21A78D17 CRC32;

Query Match 58.0%; Score 1355; DB 13; Length 384;
Best Local Similarity 69.9%; Pred. No. 3.9e-94;
Matches 270; Conservative 33; Mismatches 55; Indels 28; Gaps 5;
Qy 89 IMLFVPTLCMVVATKSVRYTEKNG-QLIYTPFTEDTPSVQRLNLSVNTLMIS 147
Db 1 IMLFIPVTLCLMVVATKSVRYTEKNG-QLIYTPFTEDTPSVQRLNLSVNTLMIS 60
Qy 148 VIVVMTIFLVLYKYRCYKEIHGWLIMSLMLFLFTYIYLGEVLKTYNVANDYPTLLT 207
Db 61 VIVVMTIFLVLYKYRCYKEIHGWLIMSLMLFLFTYIYLGEVLKTYNVANDYPTLLT 120
Qy 208 VNFAGVGVCHWKGPLVLAQAYLIMISALMALVFIKYLPEWSAWVILGALSVDLVAV 267
Db 121 IWNFGVGVCHWKGPLVLAQAYLIMISALMALVFIKYLPEWSAWVILGALSVDLVAV 180
Qy 268 LCPKGPLRLMVELTAQERNEPFIYALYSSAMVTVGMA-KLDPSSQGLALQLP----- 319
Db 181 LCPKGPLRLMVELTAQERNEPFIYALYSSAMVTVGMA-KLDPSSQGLALQLP----- 319
Qy 319 -----YDPEMEEDSYDSFG-----EPSYVEPPEPLTGYTGPGELEEEERGVL 362
Db 241 APTAQPEDGGFTFANVWQQHQLGPMGSTEDSRREIOLPSARPP--PVEDDEERGVL 298
Qy 363 GLGDFIFYSVLVGRAAATGSDGNTTTLACFVAILGLCLTLLLLAVFKKALPALPISITF 422
Db 299 GLGDFIFYSVLVGRASATGSDGNTTTLACFVAILGLCLTLLLLAVFKKALPALPISITF 358
Qy 423 GLIFVFSTDLNLRPMDPLASHQLYI 448
Db 359 GLIFVFATDLNLRPMDQLAVHQFYI 384

RESULT 5
O96340 PRELIMINARY; PRT; 272 AA.
AC O96340
DT 01-MAY-1999 (TremBLrel. 10, Created)
DT 01-MAY-1999 (TremBLrel. 10, Last sequence update)
DT 01-MAY-1999 (TremBLrel. 10, Last annotation update)
DE PRESENILIN (FRAGMENT).
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-CANTON S;
RX MEDLINE; 98331525.
RA MAREFANY G., DEL-FAVERO J., VALERO R., DE JONGHE C., WOODROW S.,
RA HENDRIKS L., VAN BROECKHOVEN C., GONZALEZ-DUARTE R.;
RT "Identification of a Drosophila presenilin homologue: evidence of
RT alternatively spliced forms";
RL J. Neurogenet. 12:41-54(1998).
DR EMBL; AF017025; AAD01611.1; -.
FT NON_TER 1
SQ SEQUENCE 272 AA; 29456 MW; 606B9A5C CRC32;

Query Match 24.7%; Score 576.5; DB 5; Length 272;
Best Local Similarity 47.8%; Pred. No. 5.7e-36;
Matches 133; Conservative 29; Mismatches 41; Indels 75; Gaps 8;

Qy 240 ALVFIKYLPEWSAWVILGALSVDLVAVLCPKGPLRLMVELTAQERNEPFIYALYSSAMV 299
Db 1 ALVFIKYLPEWSAWVILGALSVDLVAVLCPKGPLRLMVELTAQERNEPFIYALYSSAMV 60
Qy 300 WTV-----GMAKLDPPS-----QGALQLPYDPEMEEDSYDSFGPEPSY- 339
Db 61 YALVNTVTPQQSOAVTASSPSSSSNTTTRATQNSLA---SPEAAAAAGORTGN-SHPRQ 116
Qy 339 -----VFEPPLTGYGCE---ELE- 355
Db 117 NORDDCSVLATEAEAAAGFTQWSANLSERVARRIEQVSTQSGNAQORSNEYRTVTAPDQN 176
Qy 355 -----EEERGKVLGDFIFYSVLVGRAAATGSDGNTTTLACFVAILGLCLTLLLLAVFK 410
Db 177 HPDQGEERIKLGLGDFIFYSVLVGRASS--YGDWTTTACFVAILGLCLTLLLLAIWR 234
Qy 411 KALPALPISITITGLIFGFYFSTDLNLRPMDPLASHQLYI 448
Db 235 KALPALPISITITGLIFGFYFSTDLNLRPMDPLASHQLYI 272
RESULT 6
O95465 PRELIMINARY; PRT; 184 AA.
AC O95465
DT 01-MAY-1999 (TremBLrel. 10, Created)
DT 01-MAY-1999 (TremBLrel. 10, Last sequence update)
DT 01-MAY-1999 (TremBLrel. 10, Last annotation update)
DE MINILIN.
GN PSNI.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RA POWELL C.S., GEGG M.E., PALMER M.S.;
RT "Human presenilin 1 gene encodes an alternative protein-minilin.";
RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ008005; CAA07825.1; -.
SQ SEQUENCE 184 AA; 21073 MW; 5C6FBAEE CRC32;

Query Match 18.8%; Score 440; DB 4; Length 184;
Best Local Similarity 56.0%; Pred. No. 6.2e-26;
Matches 102; Conservative 15; Mismatches 43; Indels 22; Gaps 4;
Qy 24 ESPTPRSCOEGRQGEDGENTAQWRSQENEEDEEDPDYVCSGVP-----GRPPG--- 75
Db 3 ELPAPLSYFQNAQMSDNLNLSNTRVSDNRERQENRR-SLGHPEPLSNGRPQNSRQ 61
Qy 75 -----LEEELTKYGAHVIMLFPVTLCLMIVVATIKSVRYTEKNGQLIYTPFTED 127
Db 62 VVEQDEEEDLTKYGAHVIMLFPVTLCLMIVVATIKSVRYTEKNGQLIYTPFTED 121
Qy 128 TFSVQRLNLSVNTLMISVIVMTIFLVLYKYRCYKEIHGWLIMSLMLFLFTYIY 187
Db 122 TETVQORALHSILNAIMISVIVMTIFLVLYKYRCYK-----VSMRHSILSLTFLW 176
Qy 188 LG 189
Db 177 LG 178
RESULT 7
Q19737 PRELIMINARY; PRT; 406 AA.
AC Q19737; Q22692;
DT 01-NOV-1996 (TremBLrel. 01, Created)
DT 01-MAY-1999 (TremBLrel. 10, Last sequence update)
DT 01-NOV-1999 (TremBLrel. 12, Last annotation update)
DE F22E10.5 PROTEIN.
GN F22E10.5
OS Caenorhabditis elegans.

Qy	211	FGAVGVCVHWKGPLVLQQAYLIMISALMALVP	IKVLP	PEWSAWVILGALSVDVLVAVLCP	270
Db	115	IG-----	-----VILLAVATAFVGVLPMQWSEFQWGA	TVITNLISAIFY	155
Qy	271	KGPLRLVE-----	-----TAQERNEPIPPALYSISAMVTVTGMAKL	-----DPS--SQ	312
Db	156	IGP--TLAEVWVGAYDKATLRTFFAFHILP	FFIVTALAIVHLLFELHETGSNNP	SGINP	213
Qy	313	GALQLPDYDE-----	-----MEESYDSFGSEPSYEPVEFP	--PTGYGPEEL	352
Db	214	NADKIPFHPYTIKDALGFMLLSVLLLT	LFSPDSLGD---	PDNFSPLANPLTPPHIKP	270
Qy	353	EEEEERGVLGLGDFIF	-YSVLVGKAAATGSGDWN	TTLACFAVILIGLCITLLLLAVFVK	411
Db	271	E-----	-----WYFIFAVAILRSINKLGG	-----VLALLASILLLIPLLHTA	NOR 313
Qy	412	ALPALPISITFGLIIFYFSTDNLV	434		
Db	314	SMMFRIQSOT---	LFWILTANLI	333	

RESULT 11

070611	PRELIMINARY;	PRT; 1840 AA.
ID	070611	
AC	070611;	
DT	01-AUG-1998 (TREMELrel. 07, Created)	
DT	01-AUG-1998 (TREMELrel. 07, Last sequence update)	
DT	01-NOV-1999 (TREMELrel. 12, Last annotation update)	
DE	RAT SKELETAL MUSCLE TYPE 1 VOLTAGE-GATED SODIUM CHANNEL	
DE	(RSKM1) VARIANT (RSKM1).	
DN	SCN4A.	
GN		
OS	Rattus norvegicus (Rat).	
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;	
OC	Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.	
[1]		
RP	SEQUENCE FROM N.A.	
RP	STRAIN=COPENHAGEN; TISSUE=PROSTATE;	
RC	DISS J.K., STEWART D., BLACK J.A., FRASER S.P., DIBB-HAJJ S.,	
RA	WAXMAN S.G., ARCHER S., DJANGOZ M.B.A.;	
RL	FEBS Lett. 0:0-0(0).	
DR	EMBL; Y17153; CAA76659.1; -.	
DR	PFAM; PF00520; ion.trans; 4.	
DR	PFAM; PF00612; IQ_1	
DR	PRINTS; PR00170; NACHANNEL.	
DR	ionic channel.	
SK	SEQUENCE 1840 AA; 208823 MW; 1948B0C2 CRC32;	

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Query Match      4.7%; Score 109; DB 11; Length 1940;
Best Local Similarity 20.3%; Pred. No. 5.4;
Matches          63; Mismatches 130; Indels 192; Gaps 28;

QY  5  MASDSEEEVCDERTSLM-----SAESPTPRSCOEGRQGPPE-DGENTAQRQSENEE---- 55
      :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :|
Db  452 LAEDQKEK--EEFQOMLEKYYKHQEELEKAKAAQALESGEEADGPT-----HNKDCNG 503
      :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :|
QY  55  ----DGSEDDPRYVCSPGPRPGLSE-----ELTLKYGA-----KH 87
      :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :|
Db  504 SLDASGEKGP RP RSCSDASDAISDA MELEA HQKCPWYTKCAHKVLWNCCAPWVAFKH 563
      :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :|
QY  88  VIMLFV-----PVTLCMIWVATIKSVRFYTEKNGQLIYTPFTE---DPFSVGQRLN 137
      :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :|
Db  564 IIVLVMDPFDLGITIC-IVLNTLFWAMEHY-----PMTHEFDNVLSVGNLVFT 612
      :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :|
QY  138 SVLNTLMISVIVVMTITVLVLYKYRCKYTHGWLIMSSMLLFLFYIYLGEV----- 192
      :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :|
Db  613 GIETAEVVLKLIAMP-----YEV-----FOQGNWIFDSFVITLSLVELGLANVQGLSVL 662
      :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :|
QY  192  ----LKTYNVAMDYPTLLITVWVNF--AVGMVCVTHWKGPLVLOQAYLIMISALMAL-VFI 244
      :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :|
Db  663 RSPFLRVLKAKSWPTLNMLIKITIGNSVGAL-----GNLTVLVIAIIVFI FAVVGMQLFG 717
      :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :| :|
QY  245 K-----YLPWEV-----SAWV-----ILG-----AISVYD 263

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DE	CYTCHROME B LIGHT STRAND.
OS	Didelphis marsupialis.
OG	Mitochondrion.
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia;
OC	Metatheria; Didelphimorphia; Didelphidae; Didelphis.
RN	[1]
RP	SEQUENCE FROM N.A.
RA	PATTON J.L., REIS MARIA S.F., DA SILVA N.F.;
RL	J. Mammal. Evol. 3:3-29(1996).
CC	-I- CATALYTIC ACTIVITY: QH(2) + 2 FERRICYTOCHROME C = Q + 2
CC	FEROXYCYTOCHROME C.
CC	-I- COFACTOR: TWO HEME GROUPS (B562 AND B566) WHICH ARE NOT COVALENTLY
CC	BOUND TO THE PROTEIN (BY SIMILARITY).
DR	EMBL; U34665; AAA99746.1; -.
DR	PFAM; PF00032; cytochrome_b_C; 1.
DR	PFAM; PF00033; cytochrome_b_N; 1.
KW	Mitochondrion; Electron transport; Respiratory chain; Transmembrane;
KW	Heme.
SQ	SEQUENCE 382 AA; 43139 MW; 907FBCA3 CRC32;

Query Match	4.6%; Score 106.5; DB 8; Length 382;
Best Local Similarity	20.5%; Pred. No. 1.4;
Matches	79; Conservative 53; Mismatches 115; Indels 139; Gaps

QY	97 LCMIVVATIKSVFRTERKNGQLIYPFTEDPSVGQRLLNSVLNLTLMISVIVMTFL 156
Db	: :: : : :: : : :: : : :: : : :: : : :: : : :: :
	39 MCLLIQILT-----GLFLAWHYTSDT-----LTAFS 64
QY	157 VWLYKRCYCFIHGWLIM-----SSMLILFTYYTLGEVLKTYNVAMDYPTLLITVWN 210
Db	: :: : : :: : : :: : : :: : : :: : : :: : : :: :
	65 SV--AHICRDVNYGLIRNIHANGASMEFFCLFHVGVRGIYGSY-----LYKETWN 114
QY	211 FGAVGWVCIEHWGPELVQQAYLMI S A L M A L V E I K Y L P D S W A V I L G A I S V D L V A V L C P 270
Db	: :: : : :: : : :: : : :: : : :: : : :: : : :: :
	115 IG-----VILLTVATAFGVGVLPWGQMSFWGATVITNLLSAIY 155
QY	271 KGPLRMIVE-----TAQERNEPIFPALLYSSAMVYTWGMAK-----LDP 309
Db	: :: : : :: : : :: :~ : :: :~ : :: :~ : :: :~ : :: :~
	156 IG--NLIVIEWINGSFVDKATLTRFFAFHFILFIILAWVVVHLHFHTGSGNNPGLDP 213
QY	310 SSQAQLQLPYDP-----EMEDSYDSFGESPYDEVPEP--PLTGYPG 349
Db	: :: : : :: :~ : :: :~ : :: :~ : :: :~ : :: :~ : :: :~
	214 NSD---KIPHPHYTIKDILGLFLMIIILLSAMFSPDLGD---PDNETPANPLTPPH 267
QY	350 EELEEEERGVKLGLGDFIF-YSVLVGKAATGSGDWNTTLACFVAILIGLCITLLLV 408
Db	: :: :~ : :: :~ : :: :~ : :: :~ : :: :~ : :: :~ : :: :~
	268 IKPE-----WYFUYAIRLSIPNKUGG-----VLALLASILILIMP LLHTST 311
QY	409 FKALPALPISITFLGFVFSTDNLV 434
Db	: :: :~ : :: :~ : :: :~ : :: :~ : :: :~ : :: :~ : :: :~
	312 -QRSMMFRPISQT--LFWMLTANLI 333

Search completed: March 18, 2000, 22:07:43

Job time: 123 sec

Search completed: March 18, 2000, 22:07:43
Job time: 123 sec

=> e

E6 1 PRESENILIN-2 PS2S (MOUSE PS-2SHORT ISOFORM)/CN
E7 1 PRESENILIN-ASSOCIATED PROTEIN 1 (HPAP-1) (HUMAN INCYTE
CLONE 1353337)/CN
E8 1 PRESENILINASE/CN
E9 1 PRESEP-AGRI/CN
E10 1 PRESER ACE/CN
E11 1 PRESERCAR/CN
E12 1 PRESERIN M 72/CN
E13 1 PRESERIN T 72/CN
E14 1 PRESERT/CN
E15 1 PRESERVAC WETPROOF/CN
E16 1 PRESERVAL/CN
E17 1 PRESERVAL B/CN

=> s e4-6

L1 1 "PRESENILIN-2 (HUMAN ISOFORM)"/CN
1 "PRESENILIN-2 PS2CCAS (MOUSE)"/CN
1 "PRESENILIN-2 PS2S (MOUSE PS-2SHORT ISOFORM)"/CN
3 ("PRESENILIN-2 (HUMAN ISOFORM)"/CN OR "PRESENILIN-2 PS2CCAS
(MOUSE)"/CN OR "PRESENILIN-2 PS2S (MOUSE PS-2SHORT
ISOFORM)"/CN)

=> d 1-3 ide can

L1 ANSWER 1 OF 3 REGISTRY COPYRIGHT 2000 ACS
RN 251358-30-2 REGISTRY
CN **Presenilin-2 (human isoform) (9CI)** (CA INDEX NAME)
OTHER NAMES:
CN 5: PN: WO9960122 SEQID: 5 claimed protein
FS PROTEIN SEQUENCE
MF Unspecified
CI MAN
SR CA
LC STN Files: CA, CAPLUS

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
*** USE 'SQD' OR 'SQIDE' FORMATS TO DISPLAY SEQUENCE ***
1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 132:11416

L1 ANSWER 2 OF 3 REGISTRY COPYRIGHT 2000 ACS
RN 200445-64-3 REGISTRY
CN 330-448-presenilin-2 (Mus musculus isoform PS2Ccas) (9CI) (CA INDEX
NAME)
OTHER NAMES:
CN **Presenilin-2 PS2Ccas (mouse)**
FS PROTEIN SEQUENCE
MF Unspecified
CI MAN
SR CA
LC STN Files: CA, CAPLUS

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***